

(19) World Intellectual Property
Organization
International Bureau



(43) International Publication Date
27 May 2004 (27.05.2004)

PCT

(10) International Publication Number
WO 2004/044213 A1

(51) International Patent Classification⁷: **C12P 7/62,**
C08G 63/06

(21) International Application Number:
PCT/JP2003/013531

(22) International Filing Date: 23 October 2003 (23.10.2003)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
NO. 2002-310250 24 October 2002 (24.10.2002) JP
NO. 2003-356748 16 October 2003 (16.10.2003) JP

(71) Applicants (for all designated States except US): CANON KABUSHIKI KAISHA [JP/JP]; 3-30-2, Shimomaruko, Ohta-ku, Tokyo 146-8501 (JP). SUGAWA, Etsuko [JP/JP]; c/o CANON KABUSHIKI KAISHA, 3-30-2, Shimomaruko, Ohta-ku, Tokyo 146-8501 (JP).

(72) Inventors; and

(75) Inventors/Applicants (for US only): KENMOKU, Takashi [JP/JP]; c/o CANON KABUSHIKI KAISHA, 3-30-2, Shimomaruko, Ohta-ku, Tokyo 146-8501 (JP).

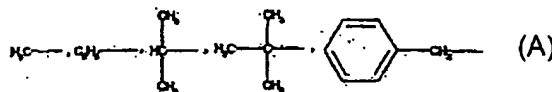
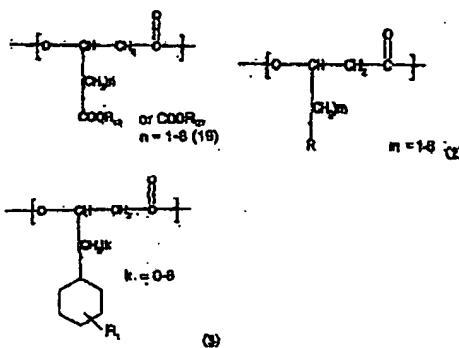
YANO, Tetsuya [JP/JP]; c/o CANON KABUSHIKI KAISHA, 3-30-2, Shimomaruko, Ohta-ku, Tokyo 146-8501 (JP). MIHARA, Chieko [JP/JP]; c/o CANON KABUSHIKI KAISHA, 3-30-2, Shimomaruko, Ohta-ku, Tokyo 146-8501 (JP). KOZAKI, Shinya [JP/JP]; c/o CANON KABUSHIKI KAISHA, 3-30-2, Shimomaruko, Ohta-ku, Tokyo 146-8501 (JP). HONMA, Tsutomu [JP/JP]; c/o CANON KABUSHIKI KAISHA, 3-30-2, Shimomaruko, Ohta-ku, Tokyo 146-8501 (JP). FUKUI, Tatsuki [JP/JP]; c/o CANON KABUSHIKI KAISHA, 3-30-2, Shimomaruko, Ohta-ku, Tokyo 146-8501 (JP). IMAMURA, Takeshi [JP/JP]; c/o CANON KABUSHIKI KAISHA, 3-30-2, Shimomaruko, Ohta-ku, Tokyo 146-8501 (JP).

(74) Agents: OKABE, Masao et al.; No. 602, Fuji Bldg., 2-3, Marunouchi 3-chome, Chiyoda-ku, Tokyo 100-0005 (JP).

(81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW.

[Continued on next page]

(54) Title: NOVEL POLYHYDROXY ALKANOATE COPOLYMER INCLUDING WITHIN MOLECULE UNIT HAVING VINYL GROUP OR CARBOXYL GROUP IN SIDE CHAIN, AND PRODUCING METHOD THEREFOR



(57) Abstract: The invention provides a PHA copolymer including at least a 3-hydroxy- ω -carboxyalkanoic acid represented by a formula (19) or (32) and simultaneously at least a unit represented by a formula (2) or a formula (3) in a molecule, a precursor PHA copolymer having a corresponding vinyl group or a corresponding alkoxycarbonyl group, a biosynthesis method thereof by microorganisms, and a method of producing a desired PHA copolymer from the precursor PHA copolymer: (wherein k, m, n are integers; R₁₈ represents H, Na, K, R₁₇ represents (A); R₁ represents a substituent on a cyclohexyl group and represents H, CN, NO₂, a halogen atom, CH₃, C₂H₅, C₃H₇, CF₃, C₂F₅, or C₃F₇; R includes a residue including a phenyl structure or a thienyl structure; these being independent for each unit).



(84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

— before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

Published:

— with international search report

- 1 -

DESCRIPTION

NOVEL POLYHYDROXY ALKANOATE COPOLYMER INCLUDING
WITHIN MOLECULE UNIT HAVING VINYL GROUP OR CARBOXYL
5 GROUP IN SIDE CHAIN, AND PRODUCING METHOD THEREFOR

TECHNICAL FIELD

The present invention relates to a polyhydroxy
alkanoate (hereinafter also abbreviated as PHA)
10 copolymer including a novel unit having a double bond
and a producing method therefor utilizing
microorganisms, also polyhydroxy alkanoate copolymer
including a novel unit having a carboxyl group or a
salt thereof, derived from the aforementioned
15 copolymer, and a producing method therefor.

Also the present invention relates to a
polyhydroxy alkanoate copolymer including a novel
unit having an ester group and a producing method
therefor utilizing microorganisms, also polyhydroxy
20 alkanoate copolymer including a novel unit having a
carboxyl group or a salt thereof, derived from the
aforementioned copolymer, and a producing method
therefor.

25 BACKGROUND ART

It has already been reported that various
microorganisms produce poly-3-hydroxybutyric acid

- 2 -

(PHB) or other poly-3-hydroxyalkanoate (PHA) and accumulate such products therein. Such PHA produced by the microorganisms can be utilized for producing various products. Also the PHA produced by
5 microorganisms, being biodegradable, has the advantage that it can be completely decomposed by the microorganisms. Therefore the PHA produced by microorganisms, when discarded, unlike the various conventional synthesized polymers, would not cause
10 pollution resulting from remaining in the natural environment. Also the PHA produced by microorganisms shows satisfactory affinity to the living tissues and is expected in the applications as the soft material for medical use.

15 However, for wider application of microorganism-produced PHA, for example for application as functional polymer, PHA having a substituent other than alkyl group in the side chain, namely "unusual PHA", is anticipated to be extremely
20 useful. Examples of hopeful substituents for this purpose include a group containing an aromatic ring (phenyl group, phenoxy group etc.), an unsaturated hydrocarbon group, an ester group, an allyl group, a cyano group, a halogenated hydrocarbon group and an
25 epoxide present on the side chain. Among these, PHA having an aromatic ring is actively investigated as follows:

- 3 -

(a) PHA containing a phenyl group or a partially substituted group thereof:

Makromol. Chem. 191, 1957-1965(1990) and Macromolecules, 24, 5256-5260(1991) report that

5 *Pseudomonas oleovorans* produces PHA containing 3-hydroxy-5-phenylvaleric acid as a unit, from 5-phenylvaleric acid as a substrate.

Macromolecules, 29, 1762-1766(1996) reports that *Pseudomonas oleovorans* produces PHA containing
10 3-hydroxy-5-(p-tolyl)valeric acid as a unit, from 5-(p-tolyl)valeric acid as a substrate.

Macromolecules, 32, 2889-2895(1999) reports that *Pseudomonas oleovorans* produces PHA containing
15 3-hydroxy-5-(2,4-dinitrophenyl)valeric acid and 3-hydroxy-5-(p-nitrophenyl)valeric acid as units, from 5-(2,4-dinitrophenyl)valeric acid as a substrate.

(b) PHA containing phenoxy group or a partially substituted group thereof:

Macromol. Chem. Phys., 195, 1665-1672(1994)
20 reports that *Pseudomonas oleovorans* produces a PHA copolymer containing 3-hydroxy-5-hydroxyvaleric acid and 3-hydroxy-9-phenoxyundecanoic acid as the units, from 11-phenoxyundecanoic acid as a substrate.

Also Japanese Patent No. 2989175 discloses
25 inventions relating to a homopolymer constituted of a 3-hydroxy-5-(monofluorophenoxy) pentanoate (3H5(MFP)P) unit or a 3-hydroxy-5-(difluorophenoxy)

- 4 -

pentanoate (3H5(DFP)P) unit, a copolymer containing either a 3H5(MFP)P unit or a 3H5(DFP)P unit or both, a novel strain of *Pseudomonas putida* capable of producing these polymers, and a method for producing the aforementioned polymers utilizing bacteria of genus *Pseudomonas*. This patent specification teaches, as the effects of such inventions, that PHA polymer having a phenoxy group substituted with 1 or 2 fluorine atoms at the end of the side chain can be biosynthesized from a long-chain fatty acid having a fluorine substituent and that thus produced PHA has a high melting point and is capable of providing stereoregularity and water repellency while maintaining satisfactory working properties.

15 In addition to the fluorine-substituted PHA having a fluorine substitution on the aromatic ring in the unit, there are also investigated PHA having a cyano group or a nitro group on the aromatic ring in the unit.

20 Can. J. Microbiol., 41, 32-43(1995) and Polymer International, 39, 205-213(1996) report production of PHA, containing 3-hydroxy-6-(p-cyanophenoxy) hexanoic acid or 3-hydroxy-6-(p-nitrophenoxy) hexanoic acid as the monomer unit, by *Pseudomonas oleovorans* ATCC 25 29347 strain and *Pseudomonas putida* KT2442 strain, from octanoic acid and 6-(p-cyanophenoxy) hexanoic acid or 6-(p-nitrophenoxy) hexanoic acid as a

- 5 -

substrate.

These references relate to PHA having an aromatic ring on the side chain, instead of alkyl groups of the usual PHA, which are effective in
5 obtaining polymer with physical properties resulting from such aromatic ring.

Also as a new category not limited to changes in the physical properties, investigations are also made for producing PHA having an appropriate
10 functional group on the side chain, thereby obtaining PHA with new functions utilizing such substituent.

As a specific method for such purpose, investigations are also made for producing PHA having, in a unit thereof, reactive group such as a bromo
15 group or a vinyl group with a high activity for example in an addition reaction to introduce an arbitrary function group in a side chain of the polymer by a chemical conversion utilizing such active group, in order to obtain PHA of multiple
20 functions.

Macromol. Rapid Commun., 20, 91-94(1999) reports production of PHA having a bromo group in a side chain by *Pseudomonas oleovorans*, and modifying the side chain with a thiolated product of acetylated
25 maltose thereby synthesizing PHA different in solubility and hydrophilicity.

Polymer, 41, 1703-1709(2000) reports producing

- 6 -

PHA, having 3-hydroxyalkenic acid with an unsaturated bond (vinyl group) at an end of a side chain as a monomer unit, by *Pseudomonas oleovorans* with 10-undecenoic acid as a substrate, followed by an
5 oxidation reaction with potassium permanganate to synthesize 3-hydroxyalkanoic acid having a diol at the end of the side chain, which PHA is reported to show such a change in solubility in solvents, as becoming soluble in polar solvents such as methanol,
10 an acetone-water (80/20, v/v) or dimethylsulfoxide and insoluble in non-polar solvents such as chloroform, tetrahydrofuran or acetone.

Also Macromolecules, 31, 1480-1486(1996) reports production of a polyester, including a unit
15 having vinyl group in a side chain by *Pseudomonas oleovorans* and epoxylating the vinyl group to obtain a polyester having an epoxy group in the side chain.

Also Polymer, 35, 2090-2097(1994) reports a crosslinking reaction within the polyester molecule
20 utilizing the vinyl group in the side chain of polyester, thereby improving physical properties of polyester.

Macromolecular chemistry, 4, 289-293(2001) reports producing PHA, including 3-hydroxy-10-
25 undecenoic acid as a monomer unit, from 10-undecenoic acid as a substrate, and then executing an oxidation reaction with potassium permanganate to obtain PHA

- 7 -

including 3-hydroxy-10-carboxydecanoic acid as a monomer unit, and reports an improvement in a decomposition thereof.

Furthermore, in order to modify physical
5 properties of PHA having an active group in a unit and to actually utilize it as a polymer, it has been studied biosynthesis of a PHA copolymer including a unit having the active group and other units; Macromolecules, 25, 1852-1857(1992) reports
10 production of a PHA copolymer including a 3-hydroxy- ω -bromoalkanoic acid unit and a linear alkanoic acid unit by *Pseudomonas oleovorans* in the presence of an ω -bromoalkanoic acid such as 11-bromoundecanoic acid, 8-bromooctanoic acid or 6-bromohexanoic acid and n-
15 nonanoic acid.

Such PHA having a highly reactive active group such as a bromo group or a vinyl group can be subjected to introduction of various functional groups or chemical modification, and such a group can
20 be a crosslinking point for a polymer, so that it is very useful means for realizing multiple functions in PHA.

Also technologies related to the present invention include a technology of oxidizing a carbon-carbon double bond with an oxidant to obtain a
25 carboxylic acid (Japanese Patent Application Laid-Open No. S59-190945, J. Chem. Soc., Perkin. Trans. 1,

- 8 -

806(1973), Org. Synth., 4, 698(1963), J. Org. Chem.,
46, 19(1981), and J. Am. Chem. Soc., 81, 4273(1959).

On the other hand, active investigations are
being made for obtaining a multi-functional PHA from
5 PHA including an ester group in a unit.

Macromol. Chem. Phys., 195, 1405-1421(1994)
reports production of a polyhydroxy alkanoate
including a unit having an ester group in a side
chain, employing *Pseudomonas oleovorans* as a
10 production microorganism and an alkanoate ester.

Also University of Massachusetts Ph. D.
Dissertation Order Number 9132875 (1991) reports
production of a polyhydroxy alkanoate including a
unit having a benzylester structure, also employing
15 *Pseudomonas oleovorans* as a production microorganism.

However, the copolymers in the foregoing
reports are comprised of a monomer unit having a
carboxyl group or an ester group at the end of a side
chain and a monomer unit having a linear alkyl group
20 (usual PHA) having a low glass transition temperature.
On the other hand, there is no report on copolymers
including 'unusual PHA having on the side chain
thereof a substituent other than a linear alkyl group,
such as a phenyl structure, a thienyl structure or a
25 cyclohexyl structure. Thus such polyhydroxy
alkanoate and a producing method therefor have been
required.

- 9 -

Also PHA having a vinyl group as an active group is a PHA copolymer with a monomer unit having a linear alkyl group (usual PHA), its low glass transition temperature and low melting point are undesirable properties in the working and the use of the polymer.

Because of the above-described situation, there have been a demand for PHA having an active group and a production method therefor, such that PHA can be produced by a microorganism at a high yield, the unit ratio of the active group can be controlled, and its physical properties can be freely regulated not to limit its application as a polymer.

15 DISCLOSURE OF THE INVENTION

As a result of intensive investigations, the present inventors have found a method of synthesizing a PHA formed by copolymerization of a unit having a vinyl group, an ester group or a carboxyl group of a high reactivity, and a unit having either one of a phenyl structure, a thienyl structure and a cyclohexyl structure which can contribute to an improvement of physical properties of the polymer, and have thus made the present invention.

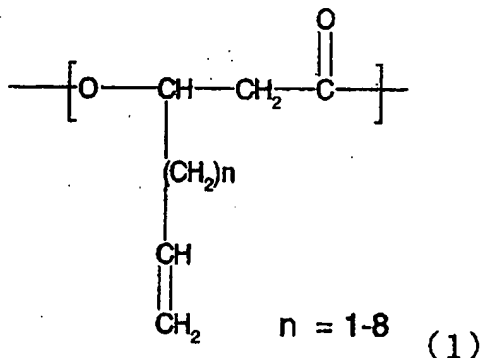
25 The present invention is outlined in the following.

[1] A polyhydroxy alkanoate copolymer including at

- 10 -

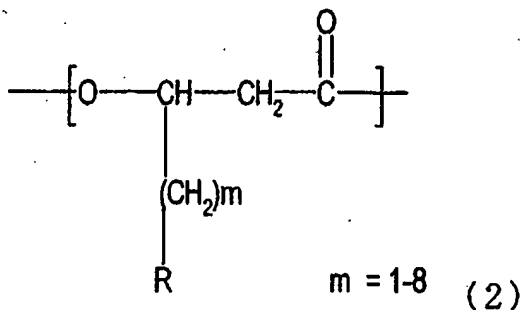
least a 3-hydroxy- ω -alkenoic acid unit represented by
 a chemical formula (1) in a molecule, and
 simultaneously at least a 3-hydroxy- ω -alkanoic acid
 unit represented by a chemical formula (2) or a 3-
 5 hydroxy- ω -cyclohexylalkanoic acid unit represented by
 a chemical formula (3) in the molecule:

[Chemical Formula (1)]



in which n represents an integer selected within a
 10 range indicated in the chemical formula; and in case
 plural units are present, n is the same or different
 for each unit;

[Chemical Formula (2)]

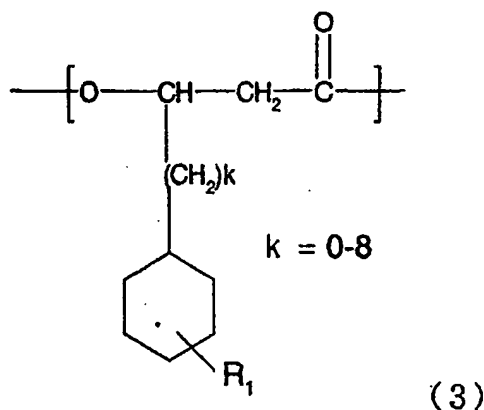


15 in which m represents an integer selected within a

- 11 -

range indicated in the chemical formula; R represents a residue having any of a phenyl structure or a thienyl structure; and in case plural units are present, m and R are the same or different for each unit;

[Chemical Formula (3)]

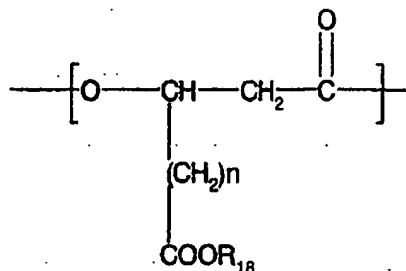


in which R_1 being a substituent on a cyclohexyl group represents a hydrogen atom, a CN group, a NO_2 group, a halogen atom, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a CF_3 group, a C_2F_5 group, or a C_3F_7 group; k represents an integer selected within a range indicated in the chemical formula; and in case plural units are present, R_1 and k may be the same or different for each unit.

[2] A polyhydroxy alkanate copolymer including at least a 3-hydroxy- ω -carboxyalkanoic acid unit represented by a chemical formula (19) or 3-hydroxy- ω -alkoxycarbonylalkanoic acid unit represented by a chemical formula (32) in a molecule, and simultaneously at least a 3-hydroxy- ω -alkanoic acid

- 12 -

unit represented by the chemical formula (2) or a 3-hydroxy- ω -cyclohexylalkanoic acid unit represented by the chemical formula (3) in the molecule,
[Chemical Formula (19)]



$n = 1-8 \quad (19)$

5

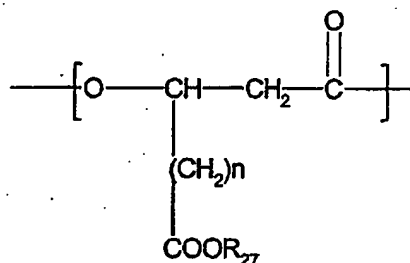
in which n represents an integer selected within a range indicated in the chemical formula; R_{18}

represents an H atom, a Na atom or a K atom:

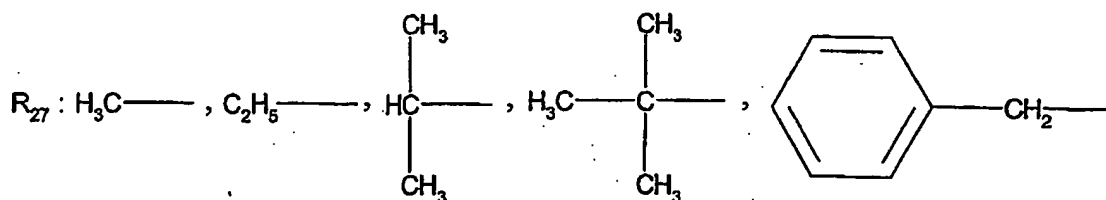
and in case plural units are present, n and R_{18} may be

10 the same or different for each unit; and

[Chemical Formula (32)]



$n = 1-8 \quad (32)$

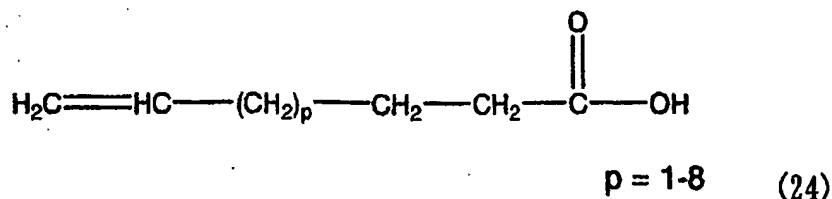


- 13 -

in which n represents an integer selected within a range indicated in the chemical formula; R_{27} represents any of residues indicated in the chemical formula; and in case plural units are present, n and R_{27} may be the same or different for each unit.

[3] A method for producing a polyhydroxy alkanoate copolymer including at least a 3-hydroxy- ω -alkenoic acid unit represented by the chemical formula (1) in a molecule, and simultaneously at least a 3-hydroxy- ω -alkanoic acid unit represented by a chemical formula (2) or a 3-hydroxy- ω -cyclohexylalkanoic acid unit represented by a chemical formula (3) in the molecule, characterized in including a biosynthesis by a production microorganism from at least an ω -alkenoic acid represented by a chemical formula (24) and at least a compound represented by a chemical formula (25) or at least an ω -cyclohexylalkanoic acid represented by a chemical formula (26) as starting materials:

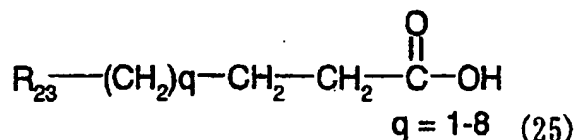
[Chemical Formula (24)]



in which p represents an integer selected within a range indicated in the chemical formula;

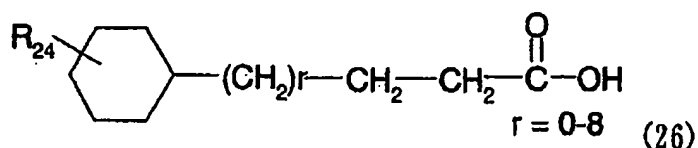
[Chemical Formula (25)]

- 14 -



in which q represents an integer selected within a range indicated in the chemical formula; and R_{23} includes a residue having a phenyl structure or a thienyl structure;

[Chemical Formula (26)]



in which R_{24} represents a substituent on a cyclohexyl group and represents an H atom, a CN group, a NO_2 group, a halogen atom, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a CF_3 group, a C_2F_5 group, or a C_3F_7 group; and r represents an integer selected within a range indicated in the chemical formula.

[4] A method for producing a polyhydroxy alkanoate copolymer including at least a 3-hydroxy- ω -carboxyalkanoic acid unit represented by the chemical formula (19) in a molecule, and simultaneously at least a 3-hydroxy- ω -alkanoic acid unit represented by the chemical formula (2) or a 3-hydroxy- ω -cyclohexylalkanoic acid unit represented by the chemical formula (3) in the molecule comprising the

- 15 -

steps of:

- preparing a polyhydroxy alkanoate copolymer including at least a 3-hydroxy- ω -alkenoic acid unit represented by the chemical formula (1) in a molecule, and simultaneously at least a 3-hydroxy- ω -alkanoic acid unit represented by the chemical formula (2) or a 3-hydroxy- ω -cyclohexylalkanoic acid unit represented by the chemical formula (3) in the molecule as a starting material, and
- oxidizing a double bond portion in the polyhydroxy alkanoate represented in the chemical formula (1) to generate the object polyhydroxy alkanoate copolymer.
- [5] A method for producing a polyhydroxy alkanoate copolymer, characterized in employing a polyhydroxy alkanoate copolymer including at least a 3-hydroxy- ω -alkoxycarbonylalkanoic acid unit represented by a chemical formula (32) in a molecule, and simultaneously at least a 3-hydroxy- ω -alkanoic acid unit represented by the chemical formula (2) or a 3-hydroxy- ω -cyclohexylalkanoic acid unit represented by the chemical formula (3) in the molecule as a starting material, and executing a hydrolysis in the presence of an acid or an alkali or executing a hydrogenolysis including a catalytic reduction, thereby generating a polyhydroxy alkanoate copolymer including at least a 3-hydroxy- ω -carboxyalkanoic acid unit represented by the chemical formula (19) in a

- 16 -

molecule, and simultaneously at least a 3-hydroxy- ω -alkanoic acid unit represented by the chemical formula (2) or a 3-hydroxy- ω -cyclohexylalkanoic acid unit represented by the chemical formula (3) in the molecule.

BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1 is a ^1H -NMR spectrum of a polyester obtained in Example 1.

Fig. 2 is a ^1H -NMR spectrum of a polyester obtained in Example 2.

Fig. 3 is a ^1H -NMR spectrum of a polyhydroxy alkanoate copolymer obtained in Example 11, and including 3-hydroxy-5-(phenylsulfanyl)valeric acid represented by a chemical formula (58), a 3-hydroxy-10-undecenoic acid represented by a chemical formula (5), 3-hydroxy-8-noneic acid represented by a chemical formula (6) and 3-hydroxy-6-heptenic acid represented by a chemical formula (7).

Fig. 4 is a ^1H -NMR spectrum of a polyhydroxy alkanoate copolymer obtained in Example 11, and including 3-hydroxy-5-(phenylsulfonyl)valeric acid represented by a chemical formula (59), a 3-hydroxy-9-carboxynonanoic acid represented by a chemical formula (54), 3-hydroxy-7-carboxyheptanoic acid represented by a chemical formula (55) and 3-hydroxy-5-carboxyvaleric acid represented by a chemical

- 17 -

formula (56).

Fig. 5 is a GC-MS TIC spectrum of a methylation decomposite of a polyester obtained in Example 34.

Fig. 6 is a mass spectrum of a peak derived
5 from a unit shown by a chemical formula (80) of a methylation decomposite of the polyester, obtained in Example 34.

Fig. 7 is a mass spectrum of a peak derived
from a unit shown by a chemical formula (81) of a
10 methylation decomposite of the polyester obtained in Example 34.

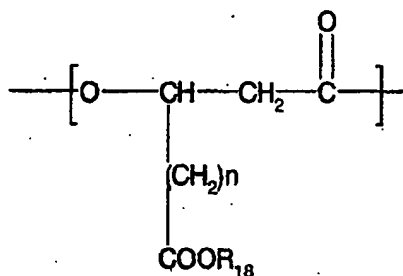
Fig. 8 is a mass spectrum of a peak derived
from a unit shown by a chemical formula (82) of a
methylation decomposite of the polyester obtained in
15 Example 34.

BEST MODE FOR CARRYING OUT THE INVENTION

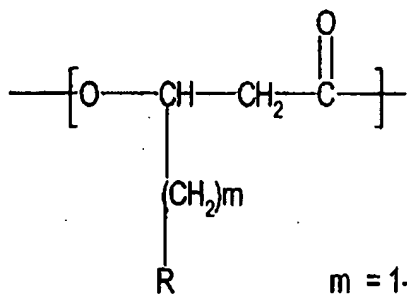
A polyhydroxy alkanoate copolymer, the final product of the present invention, is a polyhydroxy
20 alkanoate copolymer (hereinafter also called carboxyl PHA) comprising a unit having a carboxyl group on a side chain as represented by a chemical formula (19) and a unit represented by a chemical formula (2) or a chemical formula (3):

25

- 18 -

 $n = 1-8 \quad (19)$

in which n represents an integer selected within a range indicated in the chemical formula; R_{18} represents an H atom, a Na atom or a K atom;

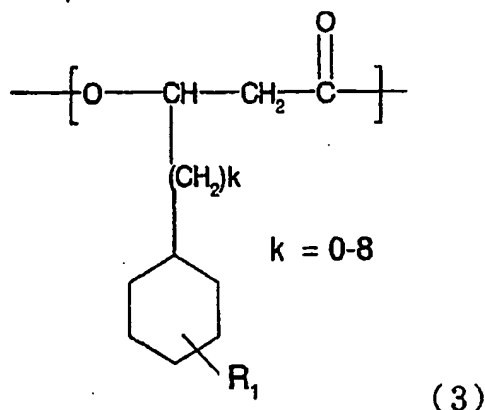
 $m = 1-8 \quad (2)$

5

in which m represents an integer selected within a range indicated in the chemical formula; R includes a residue having any of a phenyl structure or a thienyl structure; and in case plural units are present, m and R may be the same or different for each unit;

10

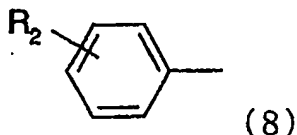
- 19 -



in which R_1 represents a substituent on a cyclohexyl group and represents a hydrogen atom, a CN group, a NO_2 group, a halogen atom, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a CF_3 group, a C_2F_5 group, or a C_3F_7 group; k represents an integer selected within a range indicated in the chemical formula; and in case plural units are present, R_1 and k may be the same or different for each unit.

10 In the present invention, R in the chemical formula (2) represents a residue having a phenyl structure or a thienyl structure selected from the group consisting of chemical formulas (8), (9), (10), (11), (12), (13), (14), (15), (16), (17) and (18):

15 the chemical formula (8):

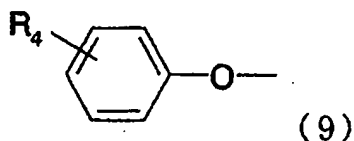


represents a group of non-substituted or substituted phenyl groups in which R_2 , a substituent on an aromatic ring and represents an H atom, represents a

halogen atom, a CN group, a NO₂ group, a CH₃ group, a C₂H₅ group, a C₃H₇ group, a CH=CH₂ group, a COOR₃ group (R₃ represents an H atom, a Na atom or a K atom) which is not included when produced by a

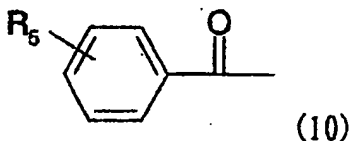
- 5 microorganism, a CF₃ group, a C₂F₅ group, or a C₃F₇ group; and in case plural units are present, R₂ is the same or different for each unit;

the chemical formula (9):



- 10 represents a group of non-substituted or substituted phenoxy groups in which R₄ represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO₂ group, a CH₃ group, a C₂H₅ group, a C₃H₇ group, a SCH₃ group, a CF₃ group, a C₂F₅ group, or a C₃F₇ group; and in case plural units are present, R₄ may be the same or different for each unit;

the chemical formula (10):

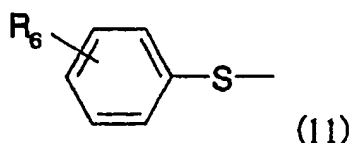


- 20 represents a group of non-substituted or substituted benzoyl groups in which R₅ represents a substituent on an aromatic ring and represents an H atom, a halogen

- 21 -

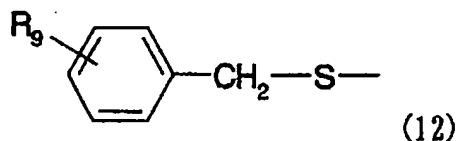
atom, a CN group, a NO₂ group, a CH₃ group, a C₂H₅ group, a C₃H₇ group, a CF₃ group, a C₂F₅ group, or a C₃F₇ group; and in case plural units are present, R₅ may be the same or different for each unit;

5 the chemical formula (11)



represents a group of substituted or non-substituted phenylsulfanyl groups in which R₆ represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO₂ group, a COOR₇ group, a SO₂R₈ group (R₇ represents either one of H, Na, K, CH₃ and C₂H₅; and R₈ represents either one of OH, ONa, OK, a halogen atom, OCH₃ and OC₂H₅), a CH₃ group, a C₂H₅ group, a C₃H₇ group, a (CH₃)₂-CH group or a (CH₃)₃-C group; and in case plural units are present, R₆ may be the same or different for each unit;

the chemical formula (12):

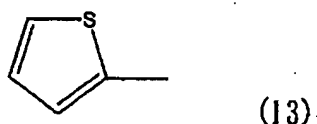


represents a group of substituted or non-substituted (phenylmethyl)sulfanyl groups in which R₉ represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO₂ group, a

- 22 -

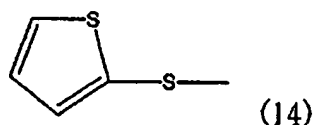
COOR₁₀ group, a SO₂R₁₁ group (R₁₀ represents either one of H, Na, K, CH₃ and C₂H₅; and R₁₁ represents either one of OH, ONa, OK, a halogen atom, OCH₃ and OC₂H₅), a CH₃ group, a C₂H₅ group, a C₃H₇ group, a (CH₃)₂-CH
 5 group or a (CH₃)₃-C group; and in case plural units are present, R₉ may be the same or different for each unit;

the chemical formula (13):



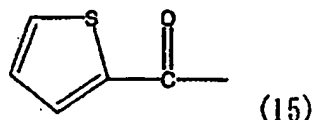
10 represents a 2-thienyl group;

the chemical formula (14)



represents a 2-thienylsulfanyl group;

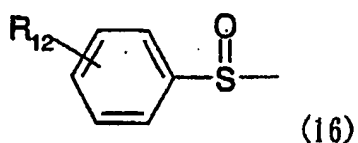
the chemical formula (15):



15

represents a 2-thienylcarbonyl group;

the chemical formula (16):

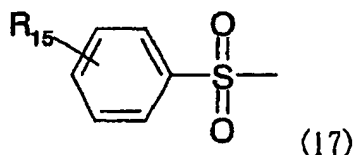


represents a group of substituted or non-substituted

- 23 -

phenylsulfinyl groups in which R_{12} represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a COOR_{13} group, a SO_2R_{14} group (R_{13} represents either one of H, Na, K, CH_3 and C_2H_5 ; and R_{14} represents either one of OH, ONa, OK, a halogen atom, OCH_3 and OC_2H_5), a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $(\text{CH}_3)_2\text{-CH}$ group or a $(\text{CH}_3)_3\text{-C}$ group; and in case plural units are present, R_{12} may be the same or different for each unit;

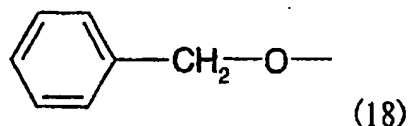
the chemical formula (17):



represents a group of substituted or non-substituted phenylsulfonyl groups in which R_{15} represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a COOR_{16} group, a SO_2R_{17} group (R_{16} represents either one of H, Na, K, CH_3 and C_2H_5 ; and R_{17} represents either one of OH, ONa, OK, a halogen atom, OCH_3 and OC_2H_5), a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $(\text{CH}_3)_2\text{-CH}$ group or a $(\text{CH}_3)_3\text{-C}$ group; and in case plural units are present, R_{15} may be the same or different for each unit; and

the chemical formula (18):

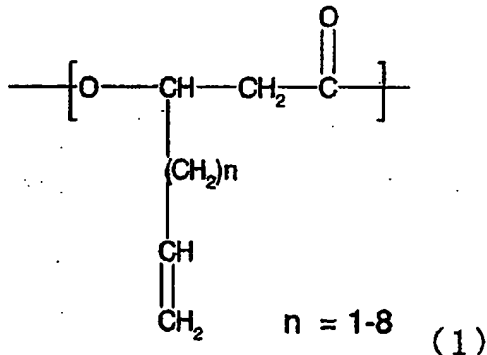
- 24 -



represents a (phenylmethyl)oxy group.

The producing methods therefor are mainly classified to:

- 5 a method of oxidizing a double bond portion in a polyhydroxy alkanooate copolymer (hereinafter also called a precursor vinyl PHA) including a 3-hydroxy- ω -alkenoic acid unit having a carbon-carbon double bond at an end of a side chain as represented in a
- 10 chemical formula (1)

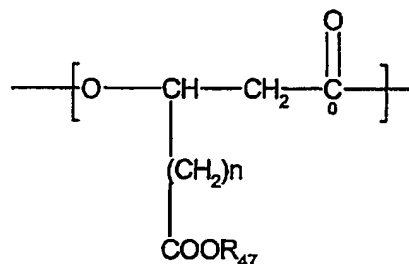


- in which n represents an integer selected within a range indicated in the chemical formula; and in case plural units are present, such units may be mutually
- 15 different), and a unit represented by a chemical formula (2) or a chemical formula (3);
- and

a method of hydrolyzing an alkoxycarbonyl portion in a polyhydroxy alkanooate copolymer

- 25 -

(hereinafter also called an alkoxycarbonyl PHA) including a 3-hydroxy- ω -alkoxyalkanoic acid unit having an ester group at an end of a side chain as represented in a chemical formula (48):



5 $n = 1-8$ (48)

in which n represents an integer selected within a range indicated in the chemical formula; R_{47} represents any of residues indicated in the chemical formula; and in case plural units are present, n and R_{41} may be the same or different for each unit, and a unit represented by a chemical formula (2) or a chemical formula (3). In the following, the precursor vinyl PHA and the precursor alkoxycarbonyl PHA may be collectively called a precursor PHA.

15 A producing method for such precursor PHA is not particularly restricted, but there can be employed a microbial production using microorganisms, a method using a genetically modified plant, or a chemical polymerization. Preferably a method by
20 microbial production is employed.

The precursor vinyl PHA and the precursor ester (alkoxycarbonyl) PHA were synthesized for the first

- 26 -

time by the present inventors, and the present invention therefore includes also the precursor vinyl PHA and the precursor ester PHA themselves, and a production process thereof by microorganisms. Also
5 such precursor vinyl PHA and precursor ester PHA can be effectively utilized not only for the carboxyl PHA which is an object of the present invention but also for introducing other functional groups.

In the following, there will be explained a
10 producing method employing each precursor PHA.

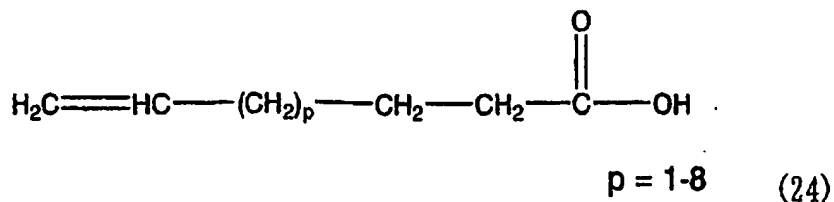
The precursor vinyl PHA can be producing by culturing a microorganism in a culture medium including an ω -alkenoic acid represented by a chemical formula (24) and a compound represented by a
15 chemical formula (25) or an ω -cyclohexylalkanoic acid represented by a chemical formula (26).

Similarly, the precursor alkoxycarbonyl PHA can be producing by culturing a microorganism in a culture medium including a carboxylic acid monoester
20 compound represented by a chemical formula (49) and a compound represented by the chemical formula (25) or an ω -cyclohexylalkanoic acid represented by the chemical formula (26).

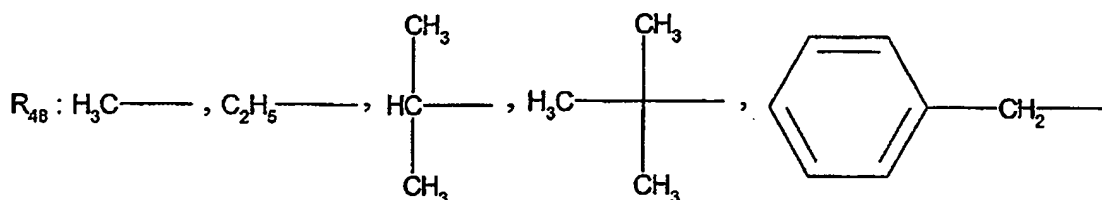
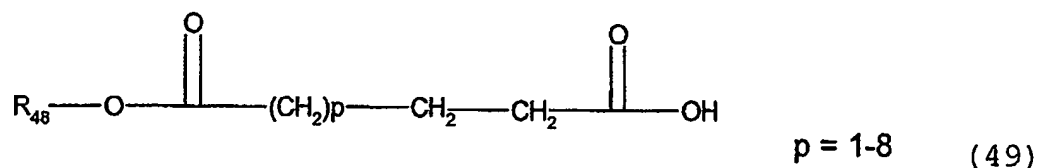
The chemical formulas (24), (49), (25) and (26)
25 are as follows:

Chemical Formula (24)

- 27 -



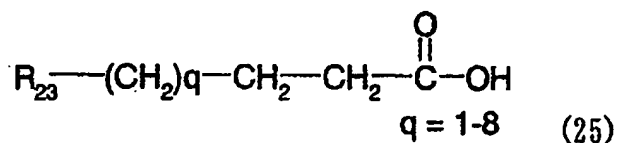
Chemical Formula (49)



5

(wherein p is an integer selected within a range indicated in the chemical formula; and R_{48} is either one of residues shown in the chemical formula.)

Chemical Formula (25)

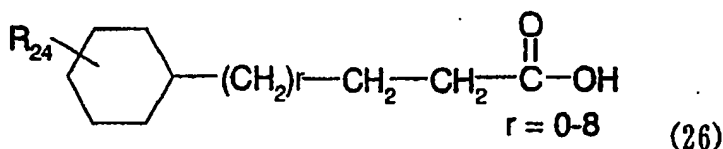


10

(wherein q is an integer selected within a range indicated in the chemical formula; and R_{23} represents a residue including a phenyl structure or a thienyl structure.)

15. Chemical Formula (26)

- 28 -



(wherein R₂₄ is a substituent on the cyclohexyl group and represents a hydrogen atom, a CN group, a NO₂ group, a halogen atom, a CH₃ group, a C₂H₅ group, a C₃H₇ group, a CF₃ group, a C₂F₅ group, or a C₃F₇ group; and r is an integer selected within a range indicated in the chemical formula.)

More specifically, each precursor PHA can be more advantageously prepared by culturing a microorganism in a culture medium containing respective raw material compounds, namely, for the precursor vinyl PHA, a combination of at least one ω-alkenoic acid represented by the chemical formula (24) and at least one compound represented by the chemical formula (25) or at least one ω-cyclohexylalkanoic acid represented by the chemical formula (26); and for the precursor alkoxycarbonyl PHA, a combination of at least one carboxylic acid monoester compound represented by the chemical formula (49) and at least one compound represented by the chemical formula (25) or at least one ω-cyclohexylalkanoic acid represented by the chemical formula (26), and further containing at least one of peptide, yeast extract, organic acid or salt thereof,

- 29 -

amino acid or a salt thereof, sugar, and linear alkanolic acid with 4 to 12 carbon atoms or salt thereof.

As preferable nutrients to be added to the culture medium, the peptide being polypeptide; one or more organic acids selected from a group of pyruvic acid, oxaloacetic acid, citric acid, isocitric acid, ketoglutaric acid, succinic acid, fumaric acid, malic acid, lactic acid and salts thereof; one or more amino acids selected from a group of glutamic acid, aspartic acid and salts thereof; and one or more sugars selected from a group of glyceraldehyde, erythrose, arabinose, xylose, glucose, galactose, mannose, fructose, glycerol, erythritol, xylitol, gluconic acid, glucuronic acid, galacturonic acid, maltose, sucrose and lactose.

In the producing method of the precursor PHA copolymer of the present invention, detailed microbial culture conditions are as follows.

The following necessary substrates and nutrients are added to an inorganic salt culture medium based on a phosphate buffer and an ammonium salt or a nitrate salt.

The raw material compound for each precursor PHA, namely, for the precursor vinyl PHA, a combination of at least an ω -alkenoic acid represented by the chemical formula (24) and at least

- 30 -

a compound represented by the chemical formula (25) or at least an ω -cyclohexylalkanoic acid represented by the chemical formula (26); or for the precursor alkoxycarbonyl PHA, a combination of at least a
5 carboxylic acid monoester compound represented by the chemical formula (49) and at least a compound represented by the chemical formula (25) or at least an ω -cyclohexylalkanoic acid represented by the chemical formula (26), is preferably contained in the
10 culture medium in a proportion of 0.01 to 1 % (w/v), further preferably 0.02 to 0.2 %.

The aforementioned nutrients as a carbon source and a nitrogen source for proliferation, and as an energy source for polyhydroxy alkanoate production
15 are preferably added to the culture medium in a proportion of 0.1 to 5 % (v/v) per medium, more preferably 0.2 to 2 %.

It can be employed any inorganic salt culture medium containing a phosphate salt and a nitrogen
20 source such as an ammonium salt or a nitrate salt, but the PHA productivity can be improved by regulating the concentration of the nitrogen source.

The culture temperature can be any temperature at which the microorganism can satisfactorily
25 proliferate, and is usually within a range of 15 to 37°C, preferably 20 to 30°C.

The culture may be carried out by any culture

- 31 -

method so long as the microorganisms can proliferate and produce PHA, such as a liquid culture or a solid culture. Also it may be batch culture, fed batch culture, semi-continuous culture or continuous
5 culture. For example, for liquid batch culture, the oxygen supply method may be shaking using a shaking flask or agitation aeration in a jar fermenter.

In order to make the microorganism produce and accumulate PHA, there can be employed, in addition to
10 the aforementioned method, a method of transferring the cell, after sufficient proliferation, to a culture medium limited in a nitrogen source such as ammonium chloride and to continue culture further in the presence of a compound being a substrate for the
15 desired unit, thereby improving the productivity.

Thus the method for producing precursor vinyl PHA of the present invention may comprise the steps of: culturing a production microorganism under the aforementioned conditions, and recovering produced
20 PHA from the cells, the PHA copolymer produced by the microorganism at least containing a 3-hydroxy- ω -alkenoic acid unit represented by the chemical formula (1), and a unit represented by the chemical formula (2) or an ω -cyclohexylalkanoic acid unit
25 represented by the chemical formula (3) in the molecule.

Also the method for producing precursor

- 32 -

alkoxycarbonyl PHA of the present invention may comprise the steps of: culturing a production microorganism under the aforementioned conditions, and recovering from the cells a polyhydroxy alkanoate copolymer produced by the microorganism which at least contains a 3-hydroxy- ω -alkoxycarbonylalkanoic acid unit represented by the chemical formula (48), and a unit represented by the chemical formula (2) or an ω -cyclohexylalkanoic acid unit represented by the chemical formula (3) in the molecule.

The object PHA can be recovered from the cells by an ordinarily employed method. For example, an extraction with an organic solvent such as chloroform, dichloromethane, ethyl acetate or acetone is most simple, but there may also be employed dioxane, tetrahydrofuran or acetonitrile. Also in a situation where an organic solvent is difficult to use, it is also possible to physically break the cells, for example by treating the cells with a surfactant such as SDS, chemicals such as hypochlorous acid and EDTA, or with an enzyme such as lysozyme, or by ultrasonic disruption, homogenizer disruption, pressure disruption, beads impulse, grinding or pounding or freeze-and-thawing, to remove cell components other than PHA and recover PHA.

A production microorganism to be employed in the production method of the present invention can be

- 33 -

any microorganisms having an ability meeting the
aforementioned conditions, but there are preferred
those belonging to the *Pseudomonas* genus, and more
preferably *Pseudomonas cichorii*, *Pseudomonas putida*,
5 *Pseudomonas fluorescense*, *Pseudomonas oleovorans*,
Pseudomonas aeruginosa, *Pseudomonas stutzeri* or
Pseudomonas jessenii. More specific examples include
Pseudomonas cichorii YN2 (FERM BP-7375), *Pseudomonas*
cichorii H45 (FERM BP-7374), *Pseudomonas jessenii*
10 P161 (FERM BP-7376), and *Pseudomonas putida* P91 (FERM
BP-7373). These four types of strains are deposited
on November 20, 2000 at International Patent Organism
Depositary, National Institute of Bioscience and
Human-Technology, Agency of Industry Science and
15 Technology (independent administrative corporation),
Tsukuba Central 6, 1-1, Higashi 1-chome, Tsukuba-shi,
Ibaraki-ken 305-8566, Japan, and described in the
Japanese Patent Application Laid-Open No. 2002-80571.

In the present invention the methods for
20 culture of the microorganism, PHA production and
accumulation by the microorganism, and for PHA
recovery from the cells are not limited to the
methods explained above.

The following is a composition of an inorganic
25 salt M9 culture medium employed in the method of the
present invention.

[M9 culture medium]

- 34 -

Na₂HPO₄ 6.3KH₂PO₄ 3.0

NaCl 0.5

NH₄Cl 1.0

5 (in g/L; pH 7.0)

For satisfactory proliferation and resulting PHA production, the above-mentioned inorganic culture medium has to be replenished with the essential trace elements by adding the following trace component

10 solution by about 0.3 % (v/v).

[Minor component solution]

nytrilotriacetic acid 1.5;

MgSO₄ 3.0;MnSO₄ 0.5;

15 NaCl 1.0;

FeSO₄ 0.1;CaCl₂ 0.1;CoCl₂ 0.1;ZnSO₄ 0.1;20 CuSO₄ 0.1;AlK(SO₄)₂ 0.1;H₃BO₃ 0.1;Na₂MoO₄ 0.1;NiCl₂ 0.1;

25 (in g/L).

The polyhydroxy alkanoates synthesized by the aforementioned producing method, a polyhydroxy

- 35 -

alkanoate copolymer including a unit represented by the chemical formula (1) and a unit represented by the chemical formula (2) or a unit represented by the chemical formula (3) can be oxidized at the carbon-carbon double bond portion to give a polyhydroxy alkanoate copolymer including a unit represented by the chemical formula (19), and a unit represented by the chemical formula (2) or a unit represented by the chemical formula (3). For obtaining a carboxylic acid by oxidizing a carbon-carbon double bond with an oxidant, there are known, for example, a method of utilizing a permanganate salt (J. Chem. Soc. Perkin. Trans. 1, 806(1973)); a method of utilizing a bichromate salt (Org. Synth., 4, 698(1963)); a method of utilizing a periodate salt (J. Org. Chem., 46, 19(1981)); a method of utilizing nitric acid (Japanese Patent Application Laid-Open No. S59-190945); a method of utilizing ozone (J. Am. Chem. Soc., 81, 4273(1959)) etc., and, on polyhydroxy alkanoate, Macromolecular chemistry, 4, 289-293(2001) reports a method of obtaining a carboxylic acid by oxidizing the carbon-carbon double bond at the end of the side chain of polyhydroxy alkanoate with potassium permanganate as an oxidant and under an acidic condition. A similar method can be utilized also in the present invention.

The oxidant to be employed in the present

- 36 -

invention, though not particularly limited, is preferably a permanganate salt. Such permanganate salt to be employed as the oxidant is usually potassium permanganate. Since the oxidation reaction is a stoichiometric reaction, the amount of the permanganate salt is usually 1 molar equivalent or more with respect to 1 mole of the unit represented by the chemical formula (1), preferably 2 to 10 molar equivalents.

For executing the reaction under an acidic condition, there is usually employed an inorganic acid such as sulfuric acid, hydrochloric acid, acetic acid or nitric acid, or an organic acid. However the use of sulfuric acid, nitric acid or hydrochloric acid may cause cleavage of an ester bond in the main chain of polyhydroxy alkanoate, thereby resulting in a decrease in the molecular weight. It is therefore preferable to employ acetic acid. An amount of acid is usually within a range of 0.2 to 2000 molar equivalents per 1 mole of the unit represented by the chemical formula (1), preferably 0.4 to 1000 molar equivalents. An amount less than 0.2 molar equivalents results in a low yield, while an amount exceeding 2000 molar equivalents generates by-products by decomposition with acid. Also a crown ether may be employed for the purpose of accelerating the reaction. In this case, crown ether and

- 37 -

permanganate salt form a complex, thereby providing an effect of increasing the reaction activity. As the crown ether, there is generally employed dibenzo-18-crown-6-ether, dicyclo-18-crown-6-ether, or 18-crown-6-ether. An amount of crown ether is generally within a range of 0.005 to 2.0 molar equivalents per 1 mole of permanganate salt, preferably 0.05 to 1.5 molar equivalents.

As a solvent to be employed in the oxidation reaction of the present invention, there may be employed any solvent inert to the reaction without particular limitation, for example water, acetone; an ether such as tetrahydrofuran or dioxane; an aromatic hydrocarbon such as benzene; an aliphatic hydrocarbon such as hexane or heptane; or a halogenated hydrocarbon such as methyl chloride, dichloromethane or chloroform. Among these solvents, in consideration of dissolving property for polyhydroxy alkanate, a halogenated hydrocarbon such as methyl chloride, dichloromethane or chloroform, or acetone is preferred.

In the aforementioned oxidation reaction of the present invention, a precursor vinyl PHA, a permanganate salt and an acid may be introduced into a solvent at a time from the beginning and reacted together, or they may be added to the reaction system one by one continuously or intermittently to be

- 38 -

- reacted. Or first a permanganate alone is dissolved or suspended in a solvent, followed by continuous or intermittent addition of a polyhydroxyalkanoate and an acid to the reaction system, or first a
- 5 polyhydroxyalkanoate alone is dissolved or suspended in a solvent, followed by continuous or intermittent addition of a permanganate and an acid to the reaction system. Further, first a polyhydroxyalkanoate and an acid are introduced into
- 10 a solvent and then a permanganate is added to the reaction system continuously or intermittently to be reacted, or first permanganate and an acid are introduced into a solvent and then polyhydroxyalkanoate is added to the reaction system
- 15 continuously or intermittently, or first a polyhydroxyalkanoate and a permanganate are introduced into a solvent and then an acid is added to the reaction system continuously and intermittently to be reacted.
- 20 A reaction temperature is selected generally within a range from -40 to 40°C, preferably -10 to 30°C. A reaction time depends on a stoichiometric ratio of the unit represented by the chemical formula (1) and permanganate salt and the reaction
- 25 temperature, but is generally selected within a range of 2 to 48 hours.

Also in the oxidation reaction of the present

- 39 -

invention, in case R_2 in the unit represented by the chemical formula (2) is a residue represented by the chemical formula (11), a sulfide bond therein may be converted into a sulfoxide or a sulfone.

5 Next, there will be explained the producing method of the precursor ester PHA of the present invention employing, as a starting material, a polyhydroxy alkanoate copolymer including a unit represented by the chemical formula (48), and a unit
10 represented by a chemical formula (2) or a unit represented by a chemical formula (3).

A precursor ester PHA synthesized can provide the carboxyl PHA by hydrolysis in the presence of an acid or an alkali or hydrogenolysis including
15 catalytic reduction of an ester bond portion shown in the chemical formula (48). Such method of hydrolysis in the presence of an acid or an alkali can be carried out by employing, in a water-miscible organic solvent such as methanol, ethanol, tetrahydrofuran,
20 dioxane, dimethylformamide or dimethylsulfoxide, an aqueous solution or an inorganic acid such as hydrochloric acid, sulfuric acid, nitric acid or phosphoric acid; an organic acid such as trifluoroacetic acid, trichloroacetic acid, p-
25 toluenesulfonic acid or methanesulfonic acid; an aqueous caustic alkali such as sodium hydroxide or potassium hydroxide; an aqueous solution or an alkali

carbonate such as sodium carbonate or potassium carbonate; or an alcoholic solution of a metal alkoxide such as sodium methoxide or sodium ethoxide. The reaction temperature is selected ordinarily from 0 to 40°C, preferably from 0 to 30°C. The reaction period is ordinarily selected from 0.5 to 48 hours. However, a hydrolysis with an acid or an alkali may also cause a cleavage of an ester bonding of the main molecular chain, thereby resulting in a decrease in the molecular weight.

Also the method of obtaining a carboxylic acid by hydrogenolysis including catalytic reduction is carried out in the following manner. Catalytic reduction is carried out in a suitable solvent and within a temperature range from -20°C to the boiling point of the used solvent, preferably from 0 to 50°C, by reacting hydrogen under a normal pressure or an elevated pressure in the presence of a reducing catalyst. Examples of the usable solvent include water, methanol, ethanol, propanol, ethyl acetate, diethyl ether, tetrahydrofuran, dioxane, benzene, toluene, dimethylformamide and pyridine. In consideration of the solubility, tetrahydrofuran, toluene or dimethylformamide is particularly preferable. As the reducing catalyst, there can be employed palladium, platinum or rhodium either singly or held on a carrier, or Raney nickel. However, the

- 41 -

catalytic reduction may also cause cleavage of an ester bonding of the main molecular chain to decrease the molecular weight.

In the following, the present invention will be explained in more details by examples thereof. These examples represent examples of the optimum embodiments of the present invention, but the present invention is by no means limited by these examples.

[Examples]

10 [Example 1]

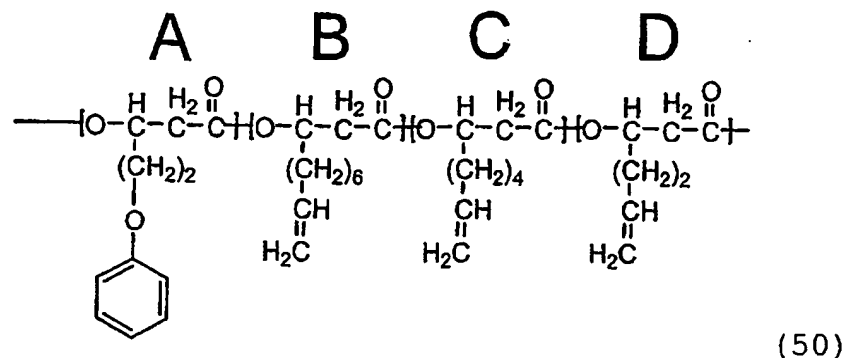
0.5% of polypeptone (supplied by Wako Pure Chemical Co.), 6 mmol/L of 5-phenoxyvaleric acid, and 1 mmol/L of 10-undecenoic acid were dissolved in 200 ml of an aforementioned M9 culture medium, which was placed in a 200 ml shaking flask, then sterilized in an autoclave and cooled to the room temperature. Then 2 ml of a culture liquid of *Pseudomonas cichorii* YN2, shake cultured in advance in an M9 culture medium containing 0.5% of polypeptone for 8 hours at 30°C, was added to the prepared culture medium, and culture was conducted for 64 hours at 30°C. After the culture, the cells were collected by centrifugation, washed with methanol and dried. The dried cells, after weighing, were put in chloroform and stirred for 72 hours at 35°C to extract a polymer. The chloroform extract was filtered, then concentrated on an evaporator, and a solid

- 42 -

precipitate formed by an addition of cold methanol was collected and dried under a reduced pressure to obtain a desired polymer.

Structure of the obtained polymer was
5 determined by ^1H -NMR (FT-NMR: Bruker DPX400; resonance frequency: 400 MHz; measured nucleus species: ^1H ; solvent: CDCl_3 ; reference: capillary-sealed TMS/ CDCl_3 ; measurement temperature: room temperature) and ^{13}C -NMR (FT-NMR: Bruker DPX400;
10 resonance frequency: 100 MHz; measured nucleus species: ^{13}C ; solvent: CDCl_3 ; reference: capillary-sealed TMS/ CDCl_3 ; measurement temperature: room temperature).

Fig. 1 shows a ^1H -NMR spectrum of the obtained
15 polymer. As a result, the obtained polymer was confirmed being a polyhydroxy alkanoate copolymer including a unit represented by the following chemical formula (50) (A : B+C+D : others (linear 3-hydroxyalkanoic acid with 4 to 12 carbon atoms and 3-
20 hydroxylalk-5-enoic acid with 10 or 12 carbon atoms) = 87 : 9 : 4). Also ^{13}C -NMR confirmed presence of the unit B which is a 3-hydroxy-10-undecenoic acid unit and both of the unit C which is a 3-hydroxy-8-nonenoic acid unit and the unit D which is a 3-
25 hydroxy-6-heptenoic acid unit, but the ratio of the units B, C and D was not determined.



The molecular weight of the obtained polymer was measured by gel permeation chromatography (GPC) (Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, molecular weight converted into polystyrene).

The obtained polymer dry weight (PDW) was 0.19 g/L and the number-averaged molecular weight was 30,000.

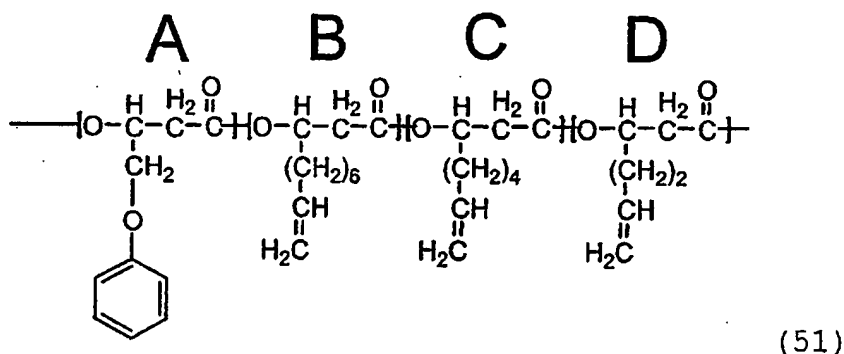
10 [Example 2]

A desired polymer was obtained in the same manner as in Example 1, except that 5-phenoxyvaleric acid employed in Example 1 was changed to 4-phenoxybutyric acid.

15 Structure of the obtained polymer was determined by $^1\text{H-NMR}$ and $^{13}\text{C-NMR}$ (FT-NMR: Bruker DPX400 as in Example 1. As a result, the obtained polymer was confirmed being a polyhydroxy alkanoate copolymer including units represented by the following chemical formula (51) (A : B+C+D : others (linear 3-hydroxyalkanoic acid with 4 to 12 carbon atoms and 3-hydroxylalk-5-enoic acid with 10 or 12

- 44 -

carbon atoms) = 72 : 11 : 15). Also ^{13}C -NMR confirmed the presence of the unit B which is a 3-hydroxy-10-undecenoic acid unit and both of the unit C which is a 3-hydroxy-8-nonenic acid unit and the unit D which is a 3-hydroxy-6-heptenoic acid unit, but the ratio of the units B, C and D was not determined.



The molecular weight of the obtained polymer was measured by GPC as in Example 1.

The obtained polymer weighed (PDW) 0.05 g/L and a number-averaged molecular weight was 25,000.

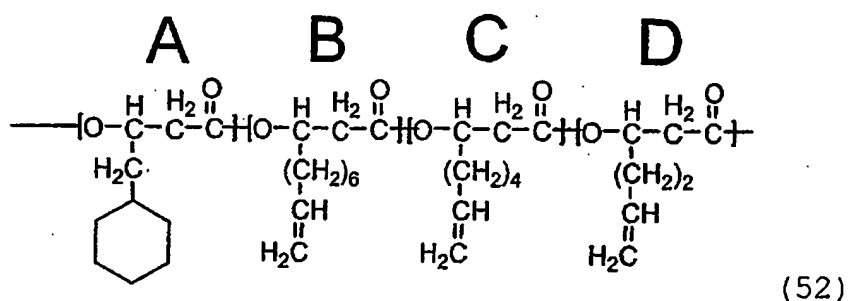
[Example 3]

A desired polymer was obtained in the same manner as in Example 1, except that 5-phenoxyvaleric acid employed in Example 1 was changed to 4-cyclohexylbutyric acid.

Structure of the obtained polymer obtained by ^1H -NMR and ^{13}C -NMR as in Example 1 was determined to confirm that the polyhydroxy alkanoate copolymer includes units represented by the following chemical formula (52) (A+others (linear 3-hydroxyalkanoic acid

- 45 -

with 4 to 12 carbon atoms and 3-hydroxylalk-5-enoic acid with 10 or 12 carbon atoms) : B+C+D = 89 : 11). Also ^{13}C -NMR confirmed the presence of the unit B being a 3-hydroxy-10-undecenoic acid unit and both of the unit C being a 3-hydroxy-8-nonenic acid unit and the unit D being a 3-hydroxy-6-heptenoic acid unit, but the ratio of the units B, C and D was not determined.



10 The molecular weight of the obtained polymer was measured by GPC as in Example 1.

The obtained polymer weighed (PDW) 0.52 g/L and the number-averaged molecular weight was 154,000.

[Example 4]

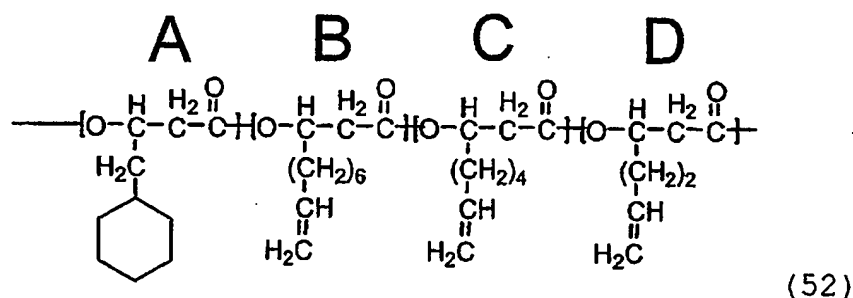
15 A desired polymer was obtained in the same manner as in Example 3, except that polypeptone employed in Example 3 was changed to yeast extract.

Structure of the obtained polymer was determined by ^1H -NMR and ^{13}C -NMR as in Example 1 to confirm the polymer being a polyhydroxy alkanoate copolymer including units represented by the following chemical formula (52) (A+others (linear 3-hydroxyalkanoic acid with 4 to 12 carbon atoms and 3-

20

- 46 -

hydroxylalk-5-enoic acid with 10 or 12 carbon atoms) : B+C+D = 85 : 15). Also ^{13}C -NMR confirmed the presence of the unit B is a 3-hydroxy-10-undecenoic acid unit and both of the unit C being a 3-hydroxy-8-nonenoic acid unit and the unit D being a 3-hydroxy-6-heptenoic acid unit, but the ratio of the units B, C and D was not determined.



The molecular weight of the obtained polymer was measured by GPC as in Example 1.

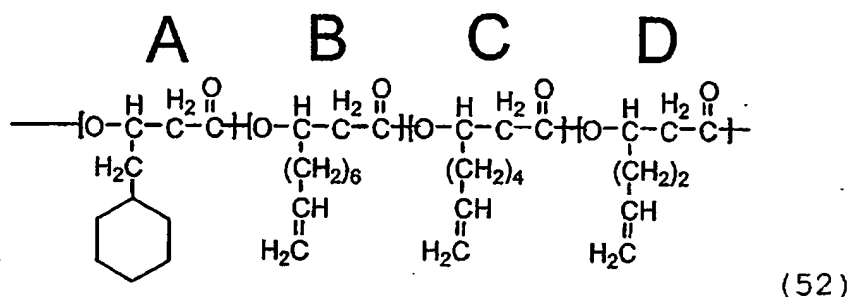
The obtained polymer weighed (PDW) 0.45 g/L and the number-averaged molecular weight was 132,000.

[Example 5]

A polymer was obtained in the same manner as in Example 3, except that the strain YN2 employed in Example 3 was replaced by *Pseudomonas cichorii* H45 and polypeptone was changed to glucose. Structure of the obtained polymer was determined by ^1H -NMR and ^{13}C -NMR as in Example 1 to confirm the polymer being a polyhydroxy alkanoate copolymer including units represented by the following chemical formula (52) (A+others (linear 3-hydroxyalkanoic acid with 4 to 12 carbon atoms and 3-hydroxylalk-5-enoic acid with 10

- 47 -

or 12 carbon atoms) : B+C+D = 83 : 17). Also ^{13}C -NMR confirmed the presence of the unit B being a 3-hydroxy-10-undecenoic acid unit and both of the unit C being a 3-hydroxy-8-nonenic acid unit and the unit D being a 3-hydroxy-6-heptenoic acid unit, but the ratio of the units B, C and D was not determined.



The molecular weight of the obtained polymer was measured by GPC as in Example 1.

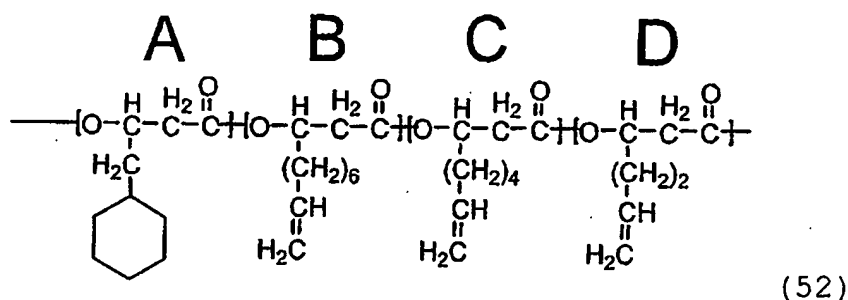
10 The obtained polymer weighed (PDW) 0.41 g/L and the number-averaged molecular weight was 164,000.

[Example 6]

A polymer was obtained in the same manner as in Example 3, except that the strain YN2 employed in Example 3 was replaced by *Pseudomonas cichorii* H45 and polypeptone was changed to sodium pyruvate. A structure determination of the obtained polymer was conducted by ^1H -NMR and ^{13}C -NMR as in Example 1 to confirm the polymer being a polyhydroxy alkanoate copolymer including units represented by the following chemical formula (52) (A+others (linear 3-hydroxyalkanoic acid with 4 to 12 carbon atoms and 3-hydroxylalk-5-enoic acid with 10 or 12 carbon

- 48 -

atoms) : B+C+D = 87 : 13). Also ^{13}C -NMR confirmed the presence of the unit B being a 3-hydroxy-10-undecenoic acid unit and the unit C being a 3-hydroxy-8-nonenic acid unit and the unit D being a 3-hydroxy-6-heptenoic acid unit, but the ratio of the units B, C and D was not determined.



The molecular weight of the obtained polymer was measured by GPC as in Example 1.

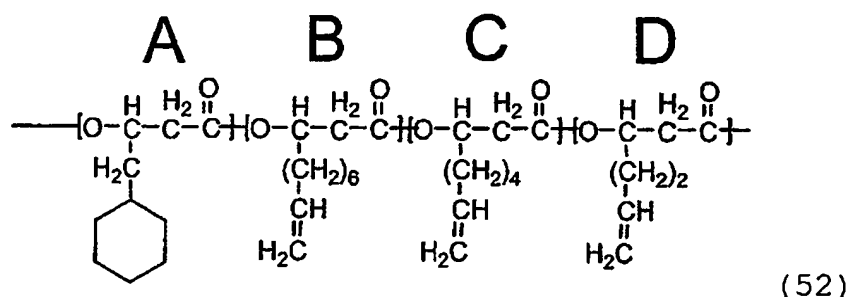
The weight of the obtained polymer (PDW) was 0.28 g/L and the number-averaged molecular weight was 156,000.

[Example 7]

A polymer was obtained in the same manner as in Example 3, except that the strain YN2 employed in Example 3 was replaced by *Pseudomonas jessenii* P161 and polypeptone was changed to sodium glutamate. Structure determination of the obtained polymer was conducted by ^1H -NMR and ^{13}C -NMR as in Example 1 to confirm the polymer being a polyhydroxy alkanoate copolymer including units represented by the following chemical formula (52) (A+others (linear 3-

hydroxyalkanoic acid with 4 to 12 carbon atoms and 3-hydroxylalk-5-enoic acid with 10 or 12 carbon atoms) : B+C+D = 88 : 12). Also ^{13}C -NMR confirmed the presence of the unit B being a 3-hydroxy-10-

5 undecenoic acid unit and both of the unit C being a 3-hydroxy-8-nonenoic acid unit and the unit D being a 3-hydroxy-6-heptenoic acid unit, but the ratio of the units B, C and D was not determined.



10 The molecular weight of the obtained polymer was measured by GPC as in Example 1.

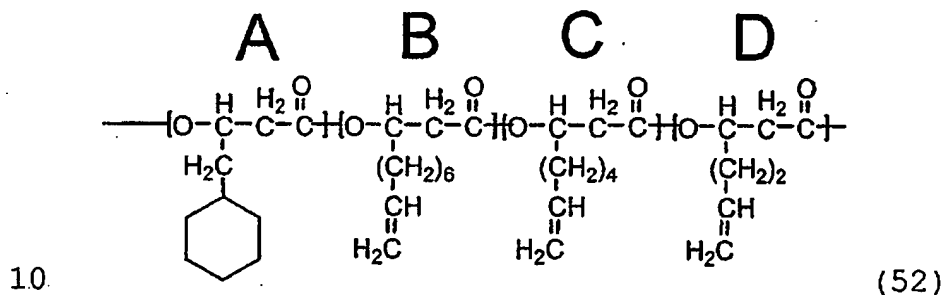
The weight of the obtained polymer (PDW) was 0.38 g/L and the number-averaged molecular weight of 145,000.

15 [Example 8]

A polymer was obtained in the same manner as in Example 3, except that the strain YN2 employed in Example 3 was replaced by *Pseudomonas jessenii* P161 and 0.5% polypeptone was changed to 0.1% of nonanic acid. The structure determination of the obtained polymer was conducted by ^1H -NMR and ^{13}C -NMR as in Example 1 to confirm the polymer being a polyhydroxy alkanoate copolymer including units represented by

- 50 -

the following chemical formula (52) (A+others (linear 3-hydroxyalkanoic acid with 4 to 12 carbon atoms and 3-hydroxylalk-5-enoic acid with 10 or 12 carbon atoms) : B+C+D = 80 : 20). Also ^{13}C -NMR confirmed the presence of the unit B being a 3-hydroxy-10-undecenoic acid unit and both of the unit C being a 3-hydroxy-8-nonenic acid unit and the unit D being a 3-hydroxy-6-heptenoic acid unit, but the ratio of the units B, C and D was not determined.



The molecular weight of the obtained polymer was measured by GPC as in Example 1.

The weight of the obtained polymer (PDW) was 0.18 g/L and the number-averaged molecular weight was 132,000.

[Example 9]

Twenty 200 ml shaking flasks were prepared, into which 0.5% of polypeptone (supplied by Wako Pure Chemical Co.), 6 mmol/L of 5-phenoxyvaleric acid, and 1 mmol/L of 10-undecenoic acid dissolved in 200 ml of an aforementioned M9 culture medium was placed, then sterilized in an autoclave and cooled to the room temperature. Then 2 ml of a culture liquid of

- 51 -

Pseudomonas cichorii YN2, shake cultured in advance in an M9 culture medium containing 0.5% of polypeptone for 8 hours at 30°C, was added to each flask, and culture was conducted for 64 hours at 30°C.

5 After the culture, all cells were collected by centrifugation, washed with methanol and dried. The dried cells, after weighing, were put in chloroform and stirred for 72 hours at 35°C to extract a polymer. The chloroform extract was filtered, then

10 concentrated on an evaporator, and a solid precipitate formed by an addition of cold methanol was collected and dried under a reduced pressure to obtain a desired polymer.

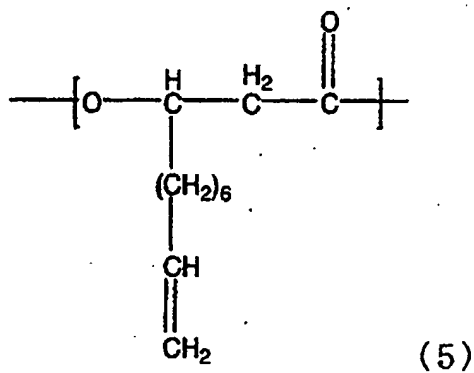
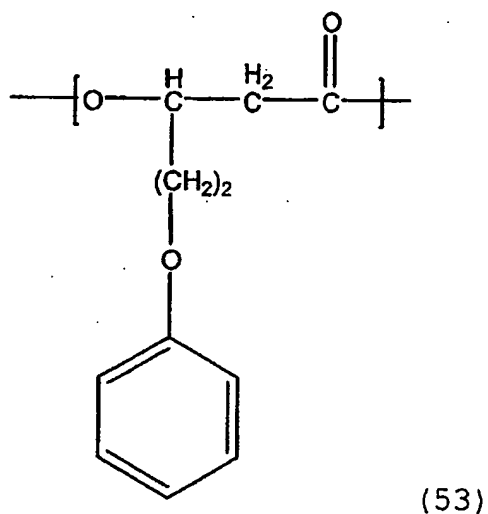
The obtained PHA polymer weighed 1528 mg (dry weight) in the present example.

The average molecular weight of the obtained PHA was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene).

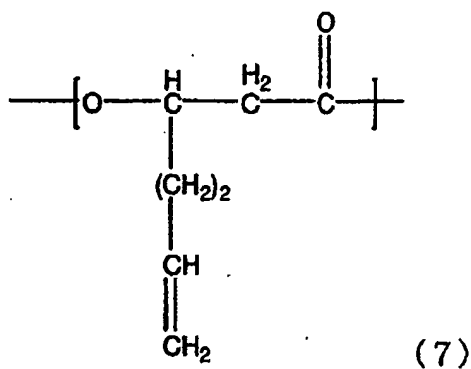
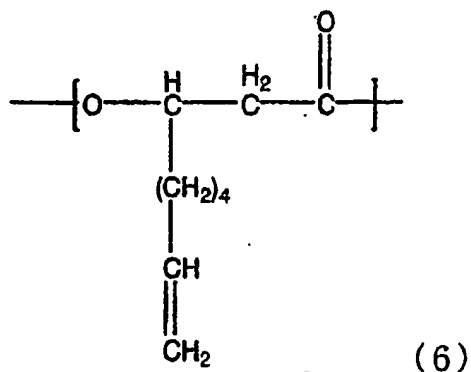
20 As a result there were obtained a number-averaged molecular weight $M_n = 104000$ and a weight-averaged molecular weight $M_w = 231000$. The structure of the obtained polymer was determined by ^1H -NMR and ^{13}C -NMR as in Example 1.

25 As a result, confirmed was a polyhydroxy alkanoate copolymer including, as monomer units, 3-hydroxy-5-phenoxyvaleric acid represented by the

following chemical formula (53), 3-hydroxy-10-undecenoic acid represented by a chemical formula (5), 3-hydroxy-8-nonenoic acid represented by a chemical formula (6) and 3-hydroxy-6-heptenoic acid represented by a chemical formula (7).



- 53 -



The proportion of such units confirmed by ¹H-NMR was: 69 mol% of 3-hydroxy-5-phenoxyvaleric acid, 23 mol% of three units of 3-hydroxy-10-undecenoic acid, 3-hydroxy-8-nonenoic acid and 3-hydroxy-6-heptenoic acid in total, and 8 mol% of others (linear 3-hydroxyalkanoic acids of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acids with 10 or 12 carbon atoms).

10 The polyhydroxy alkanoate thus obtained was utilized in the following reaction.

303 mg of polyhydroxy alkanoate were charged in a 200-ml eggplant-shaped flask and were dissolved by adding 20 ml of dichloromethane. The solution was
15 placed in an iced bath, and 3 ml of acetic acid and

- 54 -

300 mg of 18-crown-6-ether were added and agitated. Then, in an iced bath, 241 mg of potassium permanganate were slowly added and an agitation was carried out for 20 hours at the room temperature.

5 After the reaction, 50 ml of water and 500 mg of sodium bisulfite were added. Then the liquid was brought to pH = 1 by 1.0 N hydrochloric acid. After dichloromethane in the mixed solvent was distilled off in an evaporator, a polymer in the solution was

10 recovered. The polymer was recovered by washing with 100 ml of methanol and washing three times with 100 ml of purified water. A drying under a reduced pressure provided 247 mg of the desired PHA.

An average molecular weight of the obtained PHA

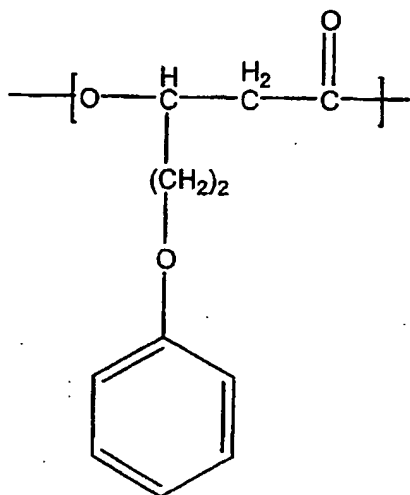
15 was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a result there were obtained a number-averaged molecular weight $M_n = 29400$ and a weight-averaged

20 molecular weight $M_w = 102800$.

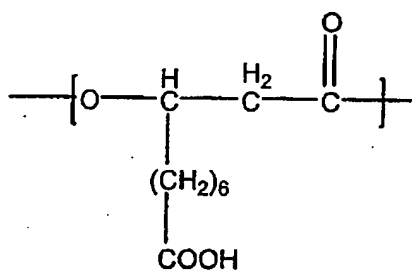
A structure determination of the obtained polymer carried out by ^1H -NMR and ^{13}C -NMR as in Example 1 confirmed a polyhydroxy alkanoate copolymer including, as monomer units, 3-hydroxy-5-

25 phenoxyvaleric acid represented by the following chemical formula (53), 3-hydroxy-9-carboxynonanoic acid represented by a chemical formula (54), 3-

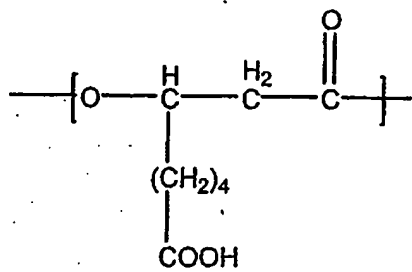
hydroxy-7-carboxyheptanoic acid represented by a chemical formula (55) and 3-hydroxy-5-carboxyvaleric acid represented by a chemical formula (56).



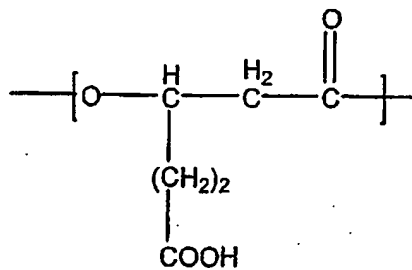
(53)



(54)



(55)



(56)

- 56 -

Also a proportion of the units of the obtained PHA was calculated by a methylesterification, utilizing trimethylsilyldiazomethane, of a carboxyl group at an end of a side chain of the PHA.

5 50 mg of the object PHA were charged in a 100-ml eggplant-shaped flask and were dissolved by adding 3.5 ml of chloroform and 0.7 ml of methanol. The solution was added with 2 ml of a 0.63 mol/L solution of trimethylsilyldiazomethane in hexane (supplied by
10 Tokyo Kasei Kogyo Co.) and was agitated for 30 minutes at the room temperature. After the reaction, the solvent was distilled off in an evaporator to recover a polymer. The polymer was recovered by washing with 50 ml of methanol. A drying under a
15 reduced pressure provided 49 mg of PHA.

NMR analysis as in Example 1 confirmed a proportion of the units in which 3-hydroxy-5-phenoxyvaleric acid was present by 83 mol%, a sum of three units of 3-hydroxy-9-carboxynonanoic acid, 3-
20 hydroxy-7-carboxyheptanoic acid and 3-hydroxy-5-carboxyvaleric acid by 8 mol%, and others (linear 3-hydroxyalkanoic acid of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or 12 carbon atoms) by 9 mol%.

25 [Example 10]

There were prepared twenty 500-ml shake flasks, and, in each, 0.5 wt.% of polypeptone (supplied by

- 57 -

Wako Pure Chemical Co.), 6 mmol/L of 4-cyclohexylbutyric acid, and 3 mmol/L of 10-undecenoic acid were dissolved in 200 ml of an aforementioned M9 culture medium, which was placed in a 500 ml shake flask, then sterilized in an autoclave and cooled to the room temperature. Then 2 ml of a culture liquid of *Pseudomonas cichorii* YN2 strain, shake cultured in advance in an M9 culture medium containing 0.5% of polypeptone for 8 hours, was added to each prepared culture medium, and culture was conducted for 60 hours at 30°C. After the culture, the culture liquids were united, and the cells were recovered by centrifuging, rinsed with methanol and dried. The dried cells, after weighing, were agitated with chloroform for 72 hours at 25°C to extract a polymer. The chloroform extract was filtered with a 0.45 µm membrane filter, then concentrated in an evaporator, and the polymer was recovered by a reprecipitation in cold methanol. A desired polymer was then obtained by drying under a reduced pressure.

According to a weighing of the obtained polymer, 1433 mg (dry weight) of PHA were obtained in the present example.

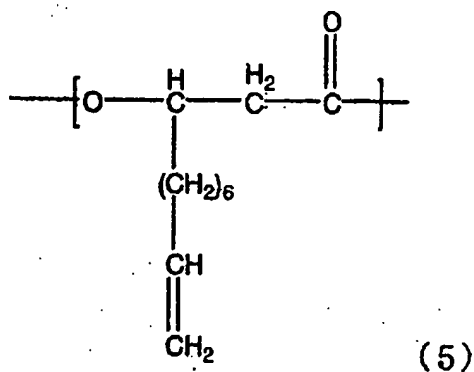
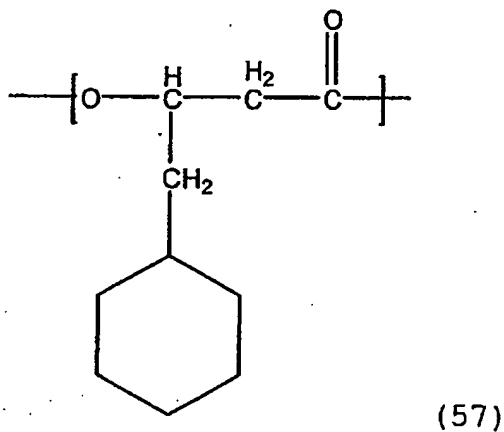
An average molecular weight of the obtained PHA was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a

- 58 -

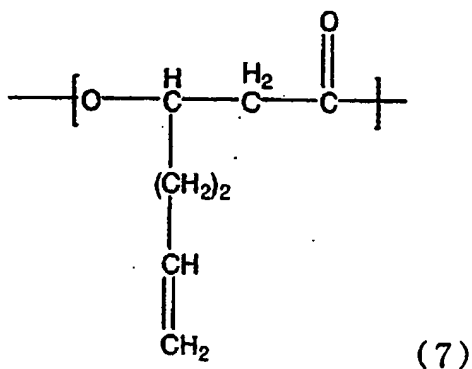
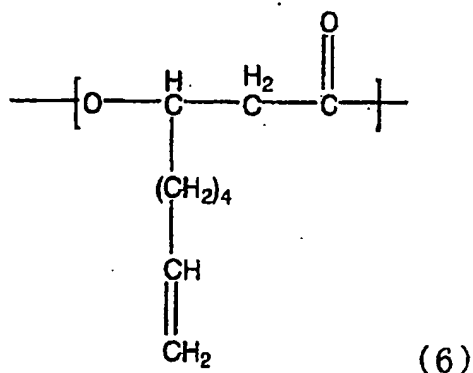
result there were obtained a number-averaged molecular weight $M_n = 143000$ and a weight-averaged molecular weight $M_w = 458000$.

A structure of the obtained PHA was determined by a NMR analysis as in Example 1.

As a result, there was confirmed a polyhydroxy alkanate copolymer including, as monomer units, 3-hydroxy-5-cyclohexylbutyric acid represented by the following chemical formula (57), 3-hydroxy-10-undecenoic acid represented by a chemical formula (5), 3-hydroxy-8-nonenoic acid represented by a chemical formula (6) and 3-hydroxy-6-heptenoic acid represented by a chemical formula (7).



- 59 -



Also a proportion of such units was confirmed by ^1H -NMR spectrum, where a sum of three units of 3-hydroxy-10-undecenoic acid, 3-hydroxy-8-nonenic acid and 3-hydroxy-6-heptenoic acid was present by 37 mol%, and 3-hydroxy-4-cyclohexylbutyric acid and others (linear 3-hydroxyalkanoic acid of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or 12 carbon atoms) by 63 mol%.

The polyhydroxy alkanoate thus obtained was utilized in the following reaction.

301 mg of polyhydroxy alkanoate were charged in a 200-ml eggplant-shaped flask and were dissolved by adding 20 ml of dichloromethane. The solution was

- 60 -

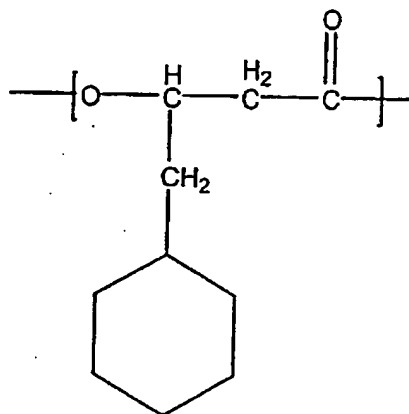
placed in an iced bath, and 3 ml of acetic acid and 541 mg of 18-crown-6-ether were added and agitated. Then, in an iced bath, 430 mg of potassium permanganate were slowly added and an agitation was carried out for 20 hours at the room temperature. After the reaction, 50 ml of water and 1000 mg of sodium bisulfite were added. Then the liquid was brought to pH = 1 by 1.0 N hydrochloric acid. After dichloromethane in the mixed solvent was distilled off in an evaporator, a polymer in the solution was recovered. The polymer was recovered by washing with 100 ml of methanol and washing three times with 100 ml of purified water. A drying under a reduced pressure provided 184 mg of the desired PHA.

15 An average molecular weight of the obtained PHA was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a result there were obtained a number-averaged
20 molecular weight $M_n = 111800$ and a weight-averaged molecular weight $M_w = 272800$.

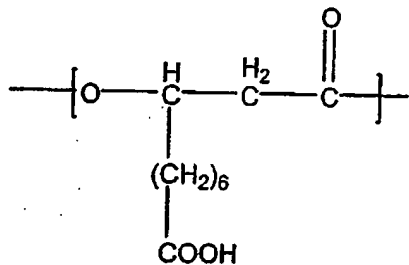
For specifying the structure of the obtained PHA, a NMR analysis was carried out under conditions same as in Example 1.

25 As a result, there was confirmed a polyhydroxy alkanoate copolymer including, as monomer units, 3-hydroxy-4-cyclohexylvaleric acid represented by the

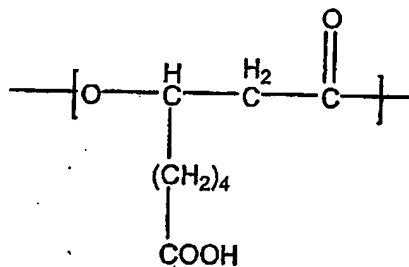
following chemical formula (57), 3-hydroxy-9-carboxynonanoic acid represented by a chemical formula (54), 3-hydroxy-7-carboxyheptanoic acid represented by a chemical formula (55) and 3-hydroxy-5-carboxyvaleric acid represented by a chemical formula (56).



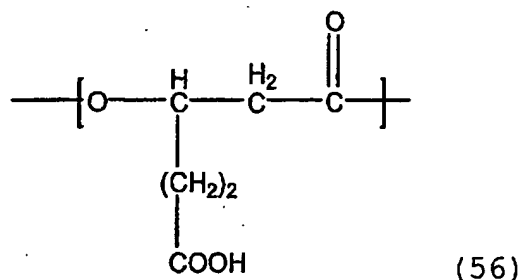
(57)



(54)



(55)



Also a proportion of the units of the obtained PHA was calculated by a methylesterification, utilizing trimethylsilyldiazomethane, of a carboxyl group at an end of a side chain of the PHA.

30 mg of the object PHA were charged in a 100-ml eggplant-shaped flask and were dissolved by adding 2.1 ml of chloroform and 0.4 ml of methanol. The solution was added with 0.9 ml of a 0.63 mol/L solution of trimethylsilyldiazomethane in hexane (supplied by Tokyo Kasei Kogyo Co.) and was agitated for 30 minutes at the room temperature. After the reaction, the solvent was distilled off in an evaporator to recover a polymer. The polymer was recovered by washing with 50 ml of methanol. A drying under a reduced pressure provided 31 mg of PHA.

A NMR analysis was carried out as in Example 1. As a result, there was confirmed a proportion of the units in which a sum of three units of 3-hydroxy-9-carboxynonanoic acid, 3-hydroxy-7-carboxyheptanoic acid and 3-hydroxy-5-carboxyvaleric acid was present by 9 mol%, and 3-hydroxy-4-cyclohexyl butyric acid and others (linear 3-hydroxyalkanoic acid of 4 to 12

- 63 -

carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or 12 carbon atoms) by 91 mol%.

[Example 11]

There were prepared three 2000-ml shake flasks, and, in each, 0.5 wt.% of polypeptone (supplied by Wako Pure Chemical Co.), 4.8 mmol/L of 5-(phenylsulfanyl)valeric acid, and 2 mmol/L of 10-undecenoic acid were dissolved in 1000 ml of an aforementioned M9 culture medium, which was placed in a 2000 ml shake flask, then sterilized in an autoclave and cooled to the room temperature. Then 10 ml of a culture liquid of *Pseudomonas cichorii* YN2 strain, shake cultured in advance in an M9 culture medium containing 0.5% of polypeptone for 8 hours, was added to each prepared culture medium, and culture was conducted for 38 hours at 30°C. After the culture, the culture liquids were united, and the cells were recovered by centrifuging, rinsed with methanol and dried. The dried cells, after weighing, were agitated with chloroform for 25 hours at 35°C to extract a polymer. The chloroform extract was filtered with a 0.45 μ m membrane filter, then concentrated in an evaporator, and the polymer was recovered by a reprecipitation in cold methanol. A desired polymer was then obtained by drying under a reduced pressure.

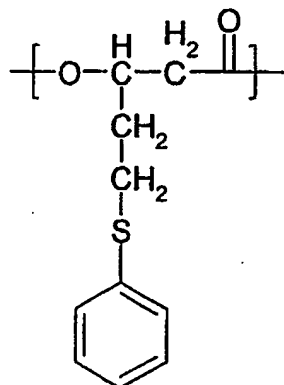
According to a weighing of the obtained polymer,

1934 mg (dry weight) of PHA were obtained in the present example.

An average molecular weight of the obtained PHA was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a result there were obtained a number-averaged molecular weight $M_n = 150000$ and a weight-averaged molecular weight $M_w = 430000$.

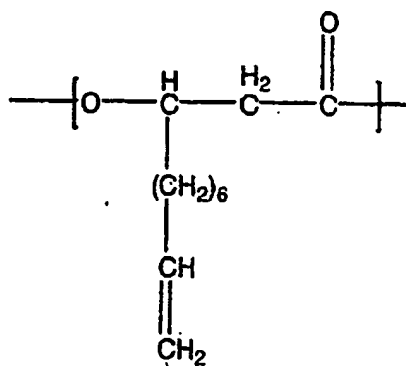
A structure of the obtained PHA was determined by a NMR analysis as in Example 1. An obtained ^1H -NMR spectrum is shown in Fig. 3.

As a result, there was confirmed a polyhydroxy alkanate copolymer including, as monomer units, 3-hydroxy-5-(phenylsulfanyl)valeric acid represented by the following chemical formula (58), 3-hydroxy-10-undecenoic acid represented by a chemical formula (5), 3-hydroxy-8-nonenoic acid represented by a chemical formula (6) and 3-hydroxy-6-heptenoic acid represented by a chemical formula (7).

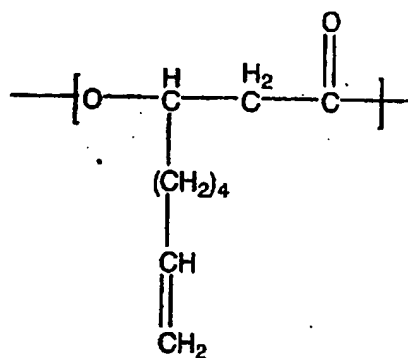


(58)

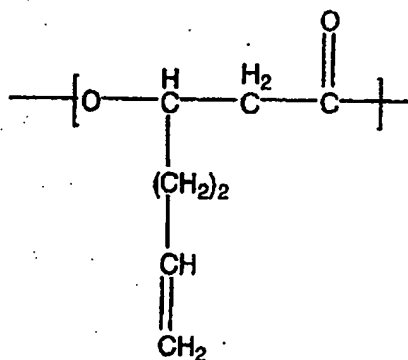
- 65 -



(5)



(6)



(7)

Also a proportion of such units was confirmed
 5 by $^1\text{H-NMR}$ spectrum, where 3-hydroxy-5-(phenylsulfanyl)valeric acid was present by 78 mol%, a sum of three units of 3-hydroxy-10-undecenoic acid, 3-hydroxy-8-nonenic acid and 3-hydroxy-6-heptenoic acid by 19 mol%, and others (linear 3-hydroxyalkanoic

- 66 -

acid of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or 12 carbon atoms) by 3 mol%.

The polyhydroxy alkanoate thus obtained was utilized in the following reaction. 302 mg of
5 polyhydroxy alkanoate were charged in a 200-ml eggplant-shaped flask and were dissolved by adding 20 ml of dichloromethane. The solution was placed in an iced bath, and 3 ml of acetic acid and 1154 mg of 18-crown-6-ether were added and agitated. Then, in an
10 iced bath, 917 mg of potassium permanganate were slowly added and an agitation was carried out for 19 hours at the room temperature. After the reaction, 50 ml of water and 3010 mg of sodium bisulfite were added. Then the liquid was brought to pH = 1 by 1.0
15 N hydrochloric acid. After dichloromethane in the mixed solvent was distilled off in an evaporator, a polymer in the solution was recovered. The polymer was recovered by washing with 100 ml of methanol and washing three times with 100 ml of purified water. A
20 drying under a reduced pressure provided 311 mg of the desired PHA.

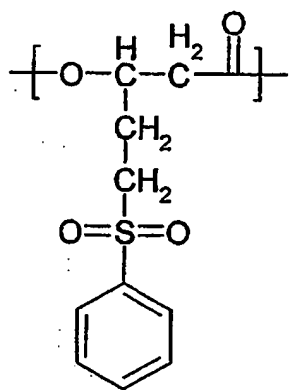
An average molecular weight of the obtained PHA was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H,
25 solvent: chloroform, converted to polystyrene). As a result there were obtained a number-averaged molecular weight $M_n = 62000$ and a weight-averaged

- 67 -

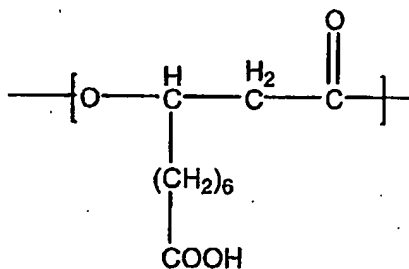
molecular weight $M_w = 260000$.

For specifying the structure of the obtained PHA, a NMR analysis was carried out under conditions same as in Example 1. An obtained ^1H -NMR spectrum is shown in Fig. 4.

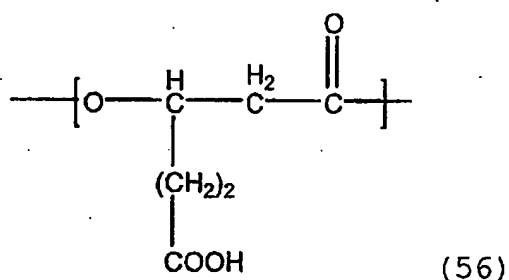
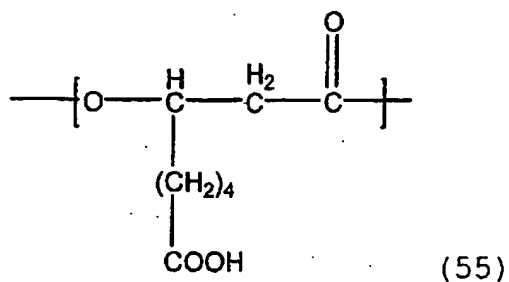
As a result, there was confirmed a polyhydroxy alkanoate copolymer including, as monomer units, 3-hydroxy-5-(phenylsulfonyl)valeric acid represented by the following chemical formula (59), 3-hydroxy-9-carboxynonanoic acid represented by a chemical formula (54), 3-hydroxy-7-carboxyheptanoic acid represented by a chemical formula (55) and 3-hydroxy-5-carboxyvaleric acid represented by a chemical formula (56).



(59)



(54)



Also a proportion of the units of the obtained
 PHA was calculated by a methylesterification,
 5 utilizing trimethylsilyldiazomethane, of a carboxyl
 group at an end of a side chain of the PHA.

30 mg of the object PHA were charged in a 100-
 ml eggplant-shaped flask and were dissolved by adding
 2.1 ml of chloroform and 0.7 ml of methanol. The
 10 solution was added with 0.5 ml of a 2 mol/L solution
 of trimethylsilyldiazomethane in hexane (supplied by
 Aldrich Inc.) and was agitated for 30 minutes at the
 room temperature. After the reaction, the solvent
 was distilled off in an evaporator to recover a
 15 polymer. The polymer was recovered by washing with
 50 ml of methanol. A drying under a reduced pressure
 provided 31 mg of PHA.

A NMR analysis was carried out as in Example 1.
 As a result, ^1H -NMR spectrum confirmed a proportion

- 69 -

of the units in which 3-hydroxy-5-(phenylsulfonyl)valeric acid was present by 89 mol%, a sum of three units of 3-hydroxy-9-carboxynonanoic acid, 3-hydroxy-7-carboxyheptanoic acid and 3-hydroxy-5-carboxyvaleric acid by 8 mol%, and others (linear 3-hydroxyalkanoic acid of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or 12 carbon atoms) by 3 mol%.

[Example 12]

There were prepared three 2000-ml shake flasks, and, in each, 0.5 wt.% of polypeptone (supplied by Wako Pure Chemical Co.), 6 mmol/L of 5-phenylvaleric acid, and 1.5 mmol/L of 10-undecenoic acid were dissolved in 1000 ml of an aforementioned M9 culture medium, which was placed in a 2000 ml shake flask, then sterilized in an autoclave and cooled to the room temperature. Then 10 ml of a culture liquid of *Pseudomonas cichorii* YN2 strain, shake cultured in advance in an M9 culture medium containing 0.5% of polypeptone for 8 hours, was added to each prepared culture medium, and culture was conducted for 60 hours at 30°C. After the culture, the culture liquids were united, and the cells were recovered by centrifuging, rinsed with methanol and dried. The dried cells, after weighing, were agitated with chloroform for 72 hours at 35°C to extract a polymer. The chloroform extract was filtered with a 0.45 µm

- 70 -

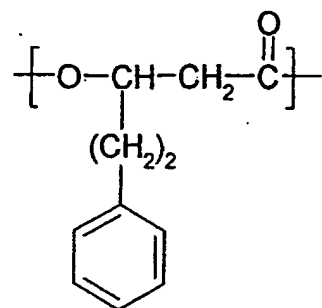
membrane filter, then concentrated in an evaporator, and the polymer was recovered by a reprecipitation in cold methanol. A desired polymer was then obtained by drying under a reduced pressure.

5 According to a weighing of the obtained polymer, 1533 mg (dry weight) of PHA were obtained in the present example.

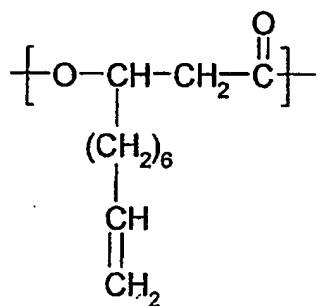
 An average molecular weight of the obtained PHA was measured by gel permeation chromatography (GPC: 10 Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a result there were obtained a number-averaged molecular weight $M_n = 72000$ and a weight-averaged molecular weight $M_w = 170000$.

15 A structure of the obtained PHA was determined by a NMR analysis as in Example 1.

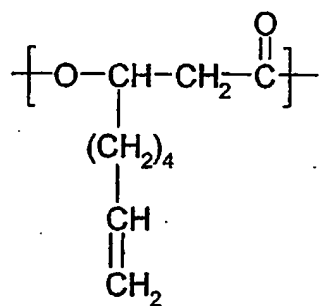
 As a result, there was confirmed a polyhydroxy alkanoate copolymer including, as monomer units, 3-hydroxy-5-phenylvaleric acid represented by the 20 following chemical formula (60), 3-hydroxy-10-undecenoic acid represented by a chemical formula (5), 3-hydroxy-8-nonenoic acid represented by a chemical formula (6) and 3-hydroxy-6-heptenoic acid represented by a chemical formula (7).



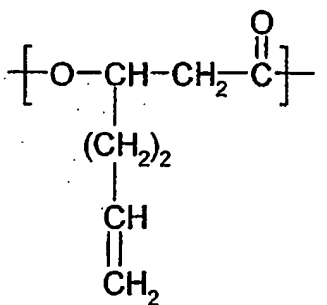
(60)



(5)



(6)



(7)

- 5 Also a proportion of such units was confirmed by ^1H -NMR spectrum, where a sum of three units of 3-hydroxy-10-undecenoic acid, 3-hydroxy-8-nonenic acid

- 72 -

and 3-hydroxy-6-heptenoic acid was present by 12 mol%,
3-hydroxy-5-phenylvaleric acid by 85 mol% and others
(linear 3-hydroxyalkanoic acid of 4 to 12 carbon
atoms and 3-hydroxyalk-5-enoic acid with 10 or 12
5 carbon atoms) by 3 mol%.

The polyhydroxy alkanoate thus obtained was
utilized in the following reaction.

1002 mg of polyhydroxy alkanoate were charged
in a 500-ml eggplant-shaped flask and were dissolved
10 by adding 60 ml of dichloromethane. The solution was
placed in an iced bath, and 10 ml of acetic acid and
537 mg of 18-crown-6-ether were added and agitated.
Then, in an iced bath, 429 mg of potassium
permanganate were slowly added and an agitation was
15 carried out for 2 hours in an iced bath and 18 hours
at the room temperature. After the reaction, 40 ml
of ethyl acetate, 30 ml of water and 1000 mg of
sodium bisulfite were added. Then the liquid was
brought to pH = 1 by 1.0 N hydrochloric acid. A
20 polymer was recovered by extraction followed by
distilling off of the solvent. The polymer was
recovered by washing with 300 ml of purified water,
then with 200 ml of methanol, three times with 200 ml
of purified water and finally with 200 ml of methanol.
25 The obtained polymer was dissolved in 10 ml of
tetrahydrofuran and dialyzed for 1 day with a
dialysis film (manufactured by Spectrum Inc.,

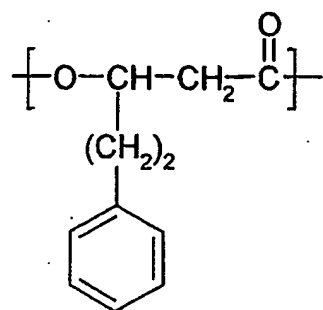
- 73 -

Spectra/Por Standard Regenerated Cellulose Dialysis Membrane 3), in a 1-L beaker containing 500 ml of methanol. The polymer present in the dialysis film was recovered and dried under a reduced pressure to
5 obtain 953 mg of a desired PHA.

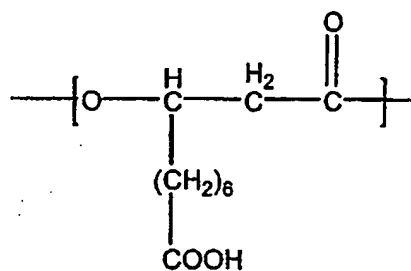
An average molecular weight of the obtained PHA was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a
10 result there were obtained a number-averaged molecular weight $M_n = 43000$ and a weight-averaged molecular weight $M_w = 94000$.

For specifying the structure of the obtained PHA, a NMR analysis was carried out under conditions
15 same as in Example 1.

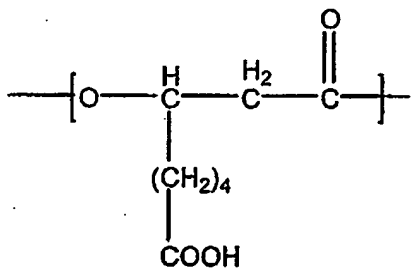
As a result, there was confirmed a polyhydroxy alkanoate copolymer including, as monomer units, 3-hydroxy-5-phenylvaleric acid represented by the following chemical formula (60), 3-hydroxy-9-
20 carboxynonanoic acid represented by a chemical formula (54), 3-hydroxy-7-carboxyheptanoic acid represented by a chemical formula (55) and 3-hydroxy-5-carboxyvaleric acid represented by a chemical formula (56).



(60)

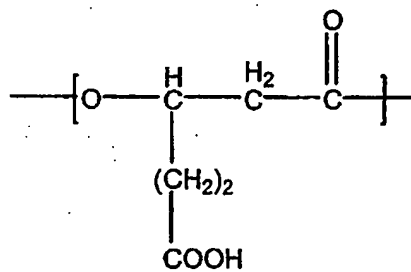


(54)



5

(55)



(56)

Also a proportion of the units of the obtained
 PHA was calculated by a methylesterification,
 10 utilizing trimethylsilyldiazomethane, of a carboxyl

- 75 -

group at an end of a side chain of the PHA.

30 mg of the object PHA were charged in a 100-ml eggplant-shaped flask and were dissolved by adding 2.1 ml of chloroform and 0.7 ml of methanol. The
5 solution was added with 0.5 ml of a 2 mol/L solution of trimethylsilyldiazomethane in hexane (supplied by Aldrich Inc.) and was agitated for 30 minutes at the room temperature. After the reaction, the solvent was distilled off in an evaporator to recover a
10 polymer. The polymer was recovered by washing with 50 ml of methanol. Drying under a reduced pressure provided 30 mg of PHA.

A NMR analysis was carried out as in Example 1. As a result, ¹H-NMR spectrum confirmed a proportion
15 of the units in which 3-hydroxy-5-phenylvaleric acid was present by 86 mol%, a sum of three units of 3-hydroxy-9-carboxynonanoic acid, 3-hydroxy-7-carboxyheptanoic acid and 3-hydroxy-5-carboxyvaleric acid by 9 mol%, and others (linear 3-hydroxyalkanoic
20 acid of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or 12 carbon atoms) by 5 mol%.

[Example 13]

500 mg of polyhydroxy alkanoate copolymer, including 3-hydroxy-5-phenylvaleric acid represented
25 by the following chemical formula (60), 3-hydroxy-10-undecenoic acid represented by a chemical formula (5), 3-hydroxy-8-nonenic acid represented by a chemical

- 76 -

formula (6) and 3-hydroxy-6-heptenoic acid represented by a chemical formula (7) as monomer units used for the bacterial production in Example 12 were changed in a 500-ml three-necked flask, and were
5 suspended by adding 150 ml of distilled water containing 50 ppm of hydrogen peroxide. Ozone was blown in with a rate of 50 mg/hr and the mixture was agitated for 3 hours at the room temperature.

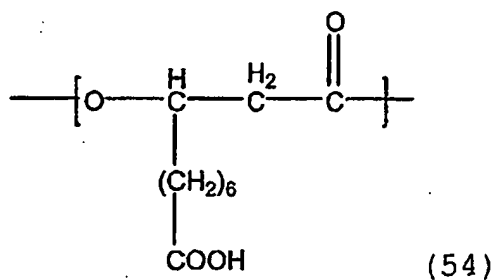
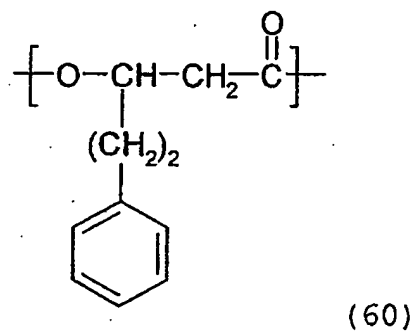
After the reaction, the reaction liquid was
10 filtered to recover a polymer. The polymer was resuspended in distilled water, and centrifuged to wash off remaining hydrogen peroxide. The obtained polymer was further dissolved in 5 ml of tetrahydrofuran and dialyzed for 1 day with a
15 dialysis film (manufactured by Spectrum Inc., Stectra/Por Standard Regenerated Cellulose Dialysis Membrane 3), in a 300-ml beaker containing 250 ml of methanol. The polymer present in the dialysis film was recovered and dried under a reduced pressure to
20 obtain 450 mg of a desired PHA.

An average molecular weight of the obtained PHA was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a
25 result there were obtained a number-averaged molecular weight $M_n = 35000$ and a weight-averaged molecular weight $M_w = 72000$.

- 77 -

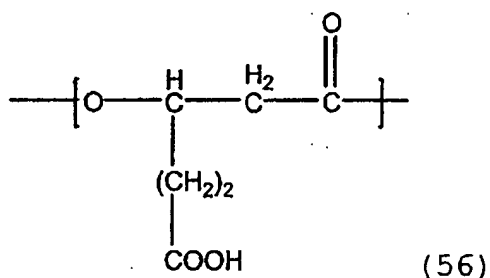
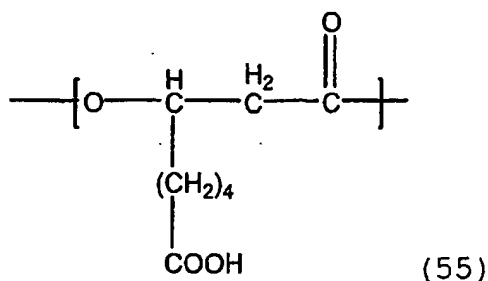
For specifying the structure of the obtained PHA, a NMR analysis was carried out under conditions same as in Example 1.

As a result, there was confirmed a polyhydroxy
 5 alkananoate copolymer including, as monomer units, 3-
 hydroxy-5-phenylvaleric acid represented by the
 following chemical formula (60), 3-hydroxy-9-
 carboxynonanoic acid represented by a chemical
 formula (54), 3-hydroxy-7-carboxyheptanoic acid
 10 represented by a chemical formula (55) and 3-hydroxy-
 5-carboxyvaleric acid represented by a chemical
 formula (56).



15

- 78 -



Also a proportion of the units of the obtained
 5 PHA was calculated by a methylesterification,
 utilizing trimethylsilyldiazomethane, of a carboxyl
 group at an end of a side chain of the PHA.

30 mg of the object PHA were charged in a 100-
 ml eggplant-shaped flask and were dissolved by adding
 10 2.1 ml of chloroform and 0.7 ml of methanol. The
 solution was added with 0.3 ml of a 2 mol/L hexane
 solution of trimethylsilyldiazomethane in hexane
 (supplied by Aldrich Inc.) and was agitated for 30
 minutes at the room temperature. After the reaction,
 15 the solvent was distilled off in an evaporator to
 recover a polymer. The polymer was recovered by
 washing with 50 ml of methanol. A drying under a
 reduced pressure provided 30 mg of PHA.

A NMR analysis was carried out as in Example 1.

- 79 -

As a result, there was confirmed a proportion of the units in which 3-hydroxy-5-phenylvaleric acid was present by 85 mol%, a sum of three units of 3-hydroxy-9-carboxynonanoic acid, 3-hydroxy-7-carboxyheptanoic acid and 3-hydroxy-5-carboxyvaleric acid by 10 mol%, and others (linear 3-hydroxyalkanoic acid of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or 12 carbon atoms) by 5 mol%.

[Example 14]

10 There were prepared five 2000-ml shake flasks, and, in each, 0.5 wt.% of polypeptone (supplied by Wako Pure Chemical Co.), 6 mmol/L of 5-(4-vinylphenyl)valeric acid, and 1 mmol/L of 10-undecenoic acid were dissolved in 1000 ml of an
15 aforementioned M9 culture medium, which was placed in a 2000 ml shake flask, then sterilized in an autoclave and cooled to the room temperature. Then 10 ml of a culture liquid of *Pseudomonas cichorii* YN2 strain, shake cultured in advance in an M9 culture
20 medium containing 0.5% of polypeptone for 8 hours, was added to each prepared culture medium, and culture was conducted for 60 hours at 30°C. After the culture, the culture liquids were united, and the cells were recovered by centrifuging, rinsed with
25 methanol and dried. The dried cells, after weighing, were agitated with chloroform for 72 hours at 25°C to extract a polymer. The chloroform extract was

- 80 -

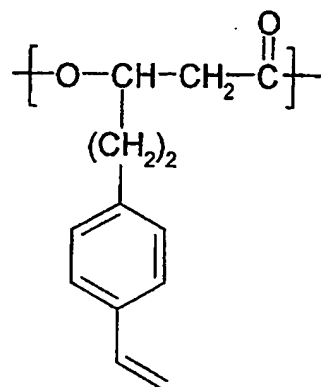
filtered with a 0.45 μ m membrane filter, then concentrated in an evaporator, and the polymer was recovered by a reprecipitation in cold methanol. A desired polymer was then obtained by drying under a
5 reduced pressure.

According to a weighing of the obtained polymer, 1097 mg (dry weight) of PHA were obtained in the present example.

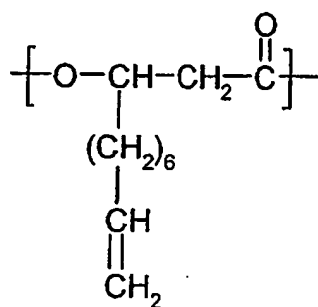
An average molecular weight of the obtained PHA
10 was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a result there were obtained a number-averaged molecular weight $M_n = 70000$ and a weight-averaged
15 molecular weight $M_w = 150000$.

A structure of the obtained PHA was determined by a NMR analysis as in Example 1.

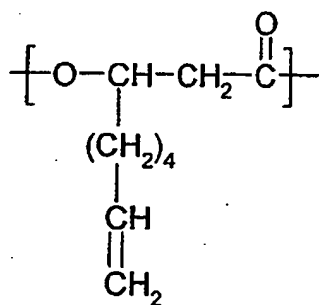
As a result, there was confirmed a polyhydroxy alkanoate copolymer including, as monomer units, 3-
20 hydroxy-5-(4-vinylphenyl)valeric acid represented by the following chemical formula (61), 3-hydroxy-10-undecenoic acid represented by a chemical formula (5), 3-hydroxy-8-nonenic acid represented by a chemical formula (6) and 3-hydroxy-6-heptenoic acid
25 represented by a chemical formula (7).



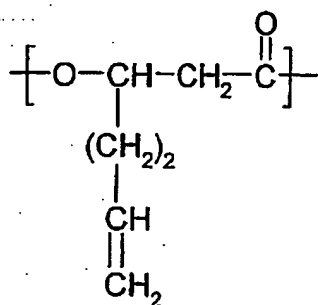
(61)



(5)



(6)



(7)

5

Also a proportion of such units was confirmed by ^1H -NMR spectrum, where a sum of three units of 3-

- 82 -

hydroxy-10-undecenoic acid, 3-hydroxy-8-nonenoic acid and 3-hydroxy-6-heptenoic acid was present by 9 mol%, 3-hydroxy-5-(4-vinylphenyl)valeric acid by 84 mol% and others (linear 3-hydroxyalkanoic acid of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or 12 carbon atoms) by 7 mol%.

[Example 15]

There were prepared twenty 2000-ml shake flasks, and, in each, 0.5 wt.% of polypeptone (supplied by Wako Pure Chemical Co.), 6 mmol/L of 5-benzoylvaleric acid, and 1 mmol/L of 10-undecenoic acid were dissolved in 1000 ml of an aforementioned M9 culture medium, which was placed in a 2000 ml shake flask, then sterilized in an autoclave and cooled to the room temperature. Then 10 ml of a culture liquid of *Pseudomonas cichorii* YN2 strain, shake cultured in advance in an M9 culture medium containing 0.5% of polypeptone for 8 hours, was added to each prepared culture medium, and culture was conducted for 60 hours at 30°C. After the culture, the culture liquids were united, and the cells were recovered by centrifuging, rinsed with methanol and dried. The dried cells, after weighing, were agitated with chloroform for 72 hours at 25°C to extract a polymer. The chloroform extract was filtered with a 0.45 µm membrane filter, then concentrated in an evaporator, and the polymer was recovered by a reprecipitation in

- 83 -

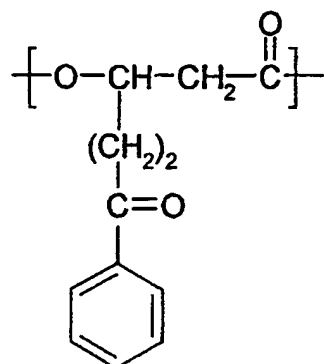
cold methanol. A desired polymer was then obtained by drying under a reduced pressure.

According to a weighing of the obtained polymer, 1027 mg (dry weight) of PHA were obtained in the
5 present example.

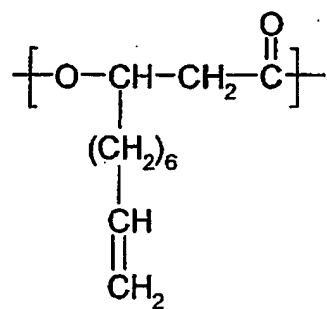
An average molecular weight of the obtained PHA was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a
10 result there were obtained a number-averaged molecular weight $M_n = 120000$ and a weight-averaged molecular weight $M_w = 370000$.

A structure of the obtained PHA was determined by a NMR analysis as in Example 1.

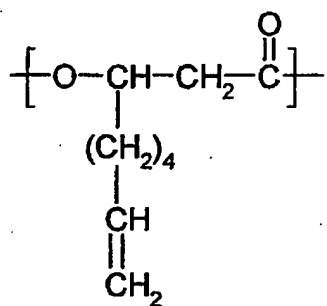
15 As a result, there was confirmed a polyhydroxy alkanoate copolymer including, as monomer units, 3-hydroxy-5-benzoylvaleric acid represented by the following chemical formula (62), 3-hydroxy-10-undecenoic acid represented by a chemical formula (5),
20 3-hydroxy-8-nonenoic acid represented by a chemical formula (6) and 3-hydroxy-6-heptenoic acid represented by a chemical formula (7).



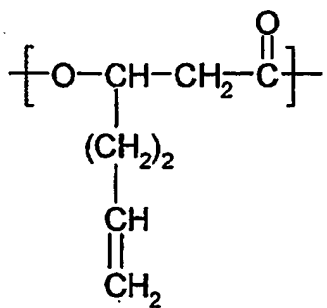
(62)



(5)



(6)



(7)

5

Also a proportion of such units was confirmed

- 85 -

by ^1H -NMR spectrum, where a sum of three units of 3-hydroxy-10-undecenoic acid, 3-hydroxy-8-nonenoic acid and 3-hydroxy-6-heptenoic acid was present by 11 mol%, 3-hydroxy-5-benzoylvaleric acid by 82 mol% and others
5 (linear 3-hydroxyalkanoic acid of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or 12 carbon atoms) by 7 mol%.

The polyhydroxy alkanoate thus obtained was utilized in the following reaction.

10 1003 mg of polyhydroxy alkanoate were charged in a 500-ml eggplant-shaped flask and were dissolved by adding 60 ml of dichloromethane. The solution was placed in an iced bath, and 10 ml of acetic acid and 410 mg of 18-crown-6-ether were added and agitated.
15 Then, in an iced bath, 327 mg of potassium permanganate were slowly added and an agitation was carried out for 2 hours in an iced bath and 18 hours at the room temperature. After the reaction, 100 ml of water and 1000 mg of sodium bisulfite were added.
20 Then the liquid was brought to pH = 1 by 1.0 N hydrochloric acid. After dichloromethane in the mixed solvent was distilled off in an evaporator, a polymer in the solution was recovered. The polymer was recovered by washing with 200 ml of purified
25 water, then with 200 ml of methanol, three times with 200 ml of purified water and finally with 200 ml of methanol. The obtained polymer was dissolved in 10

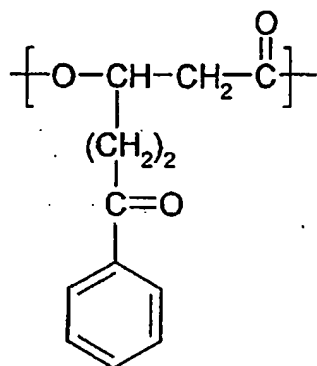
- 86 -

ml of tetrahydrofuran and dialyzed for 1 day with a dialysis film (manufactured by Spectrum Inc., Spectra/Por Standard Regenerated Cellulose Dialysis Membrane 3), in a 1-L beaker containing 500 ml of
5 methanol. The polymer present in the dialysis film was recovered and dried under a reduced pressure to obtain 948 mg of a desired PHA.

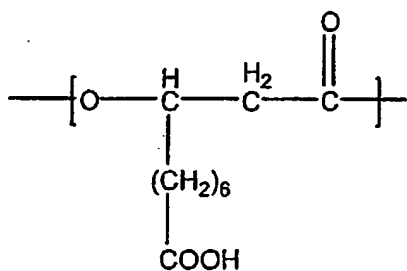
An average molecular weight of the obtained PHA was measured by gel permeation chromatography (GPC:
10 Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a result there were obtained a number-averaged molecular weight $M_n = 76000$ and a weight-averaged molecular weight $M_w = 235000$.

15 For specifying the structure of the obtained PHA, a NMR analysis was carried out under conditions same as in Example 1.

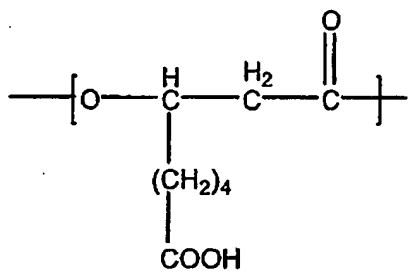
As a result, there was confirmed a polyhydroxy alkanoate copolymer including, as monomer units, 3-
20 hydroxy-5-benzoylvaleric acid represented by the following chemical formula (62), 3-hydroxy-9-carboxynonanoic acid represented by a chemical formula (54), 3-hydroxy-7-carboxyheptanoic acid represented by a chemical formula (55) and 3-hydroxy-
25 5-carboxyvaleric acid represented by a chemical formula (56).



(62)

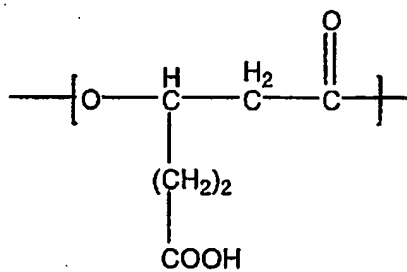


(54)



(55)

5



(56)

Also a proportion of the units of the obtained PHA was calculated by a methylesterification,

- 88 -

utilizing trimethylsilyldiazomethane, of a carboxyl group at an end of a side chain of the PHA.

30 mg of the object PHA were charged in a 100-ml eggplant-shaped flask and were dissolved by adding 2.1 ml of chloroform and 0.7 ml of methanol. The solution was added with 0.3 ml of a 2.0 mol/L hexane solution of trimethylsilyldiazomethane in hexane (supplied by Aldrich Inc.) and was agitated for 30 minutes at the room temperature. After the reaction, the solvent was distilled off in an evaporator to recover a polymer. The polymer was recovered by washing with 50 ml of methanol. A drying under a reduced pressure provided 29 mg of PHA.

A NMR analysis was carried out as in Example 1. As a result, ¹H-NMR spectrum confirmed a proportion of the units in which 3-hydroxy-5-benzoylvaleric acid was present by 84 mol%, a sum of three units of 3-hydroxy-9-carboxynonanoic acid, 3-hydroxy-7-carboxyheptanoic acid and 3-hydroxy-5-carboxyvaleric acid by 9 mol%, and others (linear 3-hydroxyalkanoic acid of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or 12 carbon atoms) by 7 mol%.

[Example 16]

There were prepared ten 2000-ml shake flasks, and, in each, 0.5 wt.% of polypeptone (supplied by Wako Pure Chemical Co.), 6 mmol/L of 5-[(phenylmethyl)sulfanyl] valeric acid, and 1.5 mmol/L

of 10-undecenoic acid were dissolved in 1000 ml of an
aforementioned M9 culture medium, which was placed in
a 2000 ml shake flask, then sterilized in an
autoclave and cooled to the room temperature. Then
5 10 ml of a culture liquid of *Pseudomonas cichorii* YN2
strain, shake cultured in advance in an M9 culture
medium containing 0.5% of polypeptone for 8 hours,
was added to each prepared culture medium, and
culture was conducted for 60 hours at 30°C. After
10 the culture, the culture liquids were united, and the
cells were recovered by centrifuging, rinsed with
methanol and dried. The dried cells, after weighing,
were agitated with chloroform for 72 hours at 25°C to
extract a polymer. The chloroform extract was
15 filtered with a 0.45 µm membrane filter, then
concentrated in an evaporator, and the polymer was
recovered by a reprecipitation in cold methanol. A
desired polymer was then obtained by drying under a
reduced pressure.

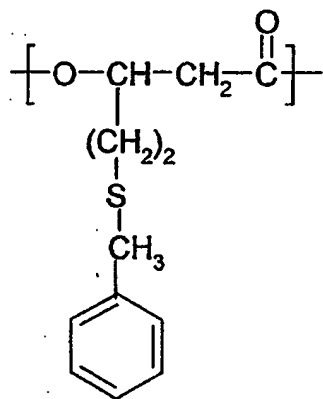
20 According to a weighing of the obtained polymer,
1714 mg (dry weight) of PHA were obtained in the
present example.

An average molecular weight of the obtained PHA
was measured by gel permeation chromatography (GPC:
25 Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H,
solvent: chloroform, converted to polystyrene). As a
result there were obtained a number-averaged

molecular weight $M_n = 110000$ and a weight-averaged molecular weight $M_w = 380000$.

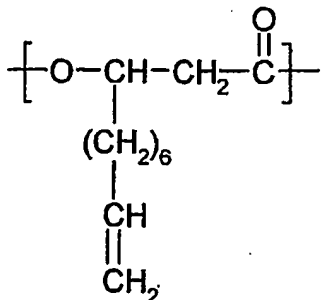
For specifying the structure of the obtained PHA, a NMR analysis was conducted under conditions similar to those in Example 1.

As a result, there was confirmed a polyhydroxy alkanate copolymer including, as monomer units, 3-hydroxy-5-[(phenylmethyl)sulfanyl]valeric acid represented by the following chemical formula (63),
 3-hydroxy-10-undecenoic acid represented by a chemical formula (5), 3-hydroxy-8-nonenoic acid represented by a chemical formula (6) and 3-hydroxy-6-heptenoic acid represented by a chemical formula (7).

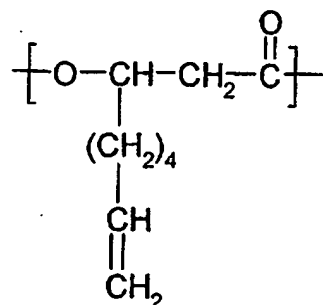


15

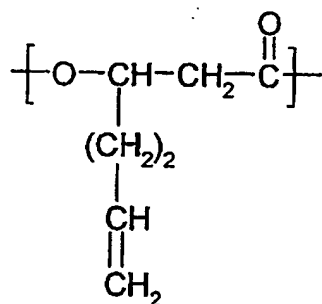
(63)



(5)



(6)



(7)

Also a proportion of such units was confirmed by ¹H-NMR spectrum, where a sum of three units of 3-hydroxy-10-undecenoic acid, 3-hydroxy-8-nonenoic acid and 3-hydroxy-6-heptenoic acid was present by 12 mol%, 3-hydroxy-5-[(phenylmethyl)sulfanyl]valeric acid by 80 mol% and others (linear 3-hydroxyalkanoic acid of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or 12 carbon atoms) by 8 mol%.

[Example 17]

There were prepared three 2000-ml shake flasks, and, in each, 0.5 wt.% of polypeptone (supplied by Wako Pure Chemical Co.), 6 mmol/L of 5-(2-thienyl)valeric acid, and 1.5 mmol/L of 10-undecenoic acid were dissolved in 1000 ml of an aforementioned M9 culture medium, which was placed in a 2000 ml

- 92 -

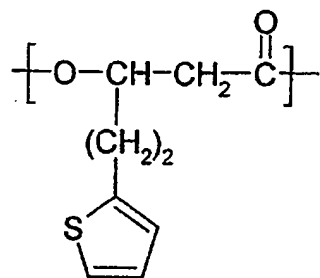
shake flask, then sterilized in an autoclave and cooled to the room temperature. Then 10 ml of a culture liquid of *Pseudomonas cichorii* YN2 strain, shake cultured in advance in an M9 culture medium
5 containing 0.5% of polypeptone for 8 hours, was added to each prepared culture medium, and culture was conducted for 60 hours at 30°C. After the culture, the culture liquids were united, and the cells were recovered by centrifuging, rinsed with methanol and
10 dried. The dried cells, after weighing, were agitated with chloroform for 72 hours at 25°C to extract a polymer. The chloroform extract was filtered with a 0.45 μ m membrane filter, then concentrated in an evaporator, and the polymer was
15 recovered by a reprecipitation in cold methanol. A desired polymer was then obtained by drying under a reduced pressure.

According to a weighing of the obtained polymer, 1171 mg (dry weight) of PHA were obtained in the
20 present example.

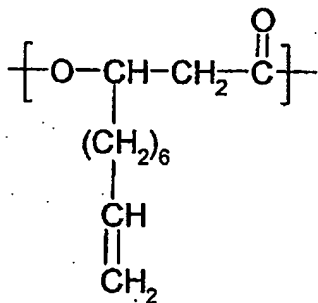
An average molecular weight of the obtained PHA was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a
25 result there were obtained a number-averaged molecular weight $M_n = 74000$ and a weight-averaged molecular weight $M_w = 180000$.

For specifying the structure of the obtained PHA, a NMR analysis was conducted under conditions similar to those in Example 1.

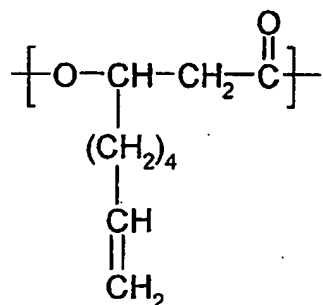
As a result, there was confirmed a polyhydroxy
 5 alkanooate copolymer including, as monomer units, 3-
 hydroxy-5-(2-thienyl)valeric acid represented by the
 following chemical formula (64), 3-hydroxy-10-
 undecenoic acid represented by a chemical formula (5),
 3-hydroxy-8-nonenoic acid represented by a chemical
 10 formula (6) and 3-hydroxy-6-heptenoic acid
 represented by a chemical formula (7).



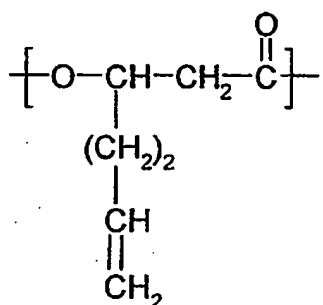
(64)



(5)



(6)



(7)

Also a proportion of such units was confirmed by ^1H -NMR spectrum, where a sum of three units of 3-hydroxy-10-undecenoic acid, 3-hydroxy-8-nonenoic acid and 3-hydroxy-6-heptenoic acid was present by 12 mol%, 3-hydroxy-5-(2-thienyl)valeric acid by 85 mol% and others (linear 3-hydroxyalkanoic acid of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or 12 carbon atoms) by 3 mol%.

The polyhydroxy alkanoate thus obtained was utilized in the following reaction.

1001 mg of polyhydroxy alkanoate were charged in a 500-ml eggplant-shaped flask and were dissolved by adding 60 ml of dichloromethane. The solution was placed in an iced bath, and 10 ml of acetic acid and 527 mg of 18-crown-6-ether were added and agitated.

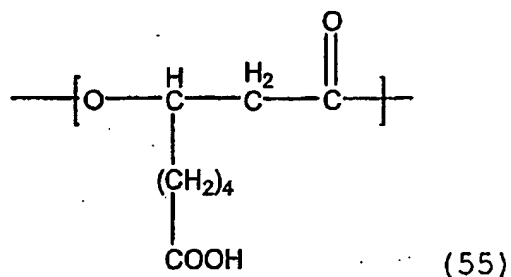
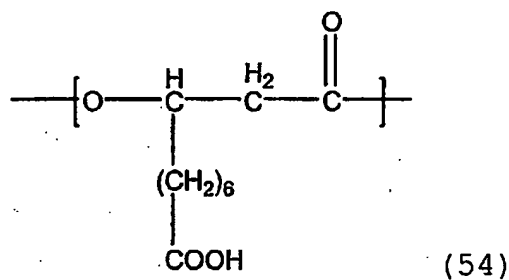
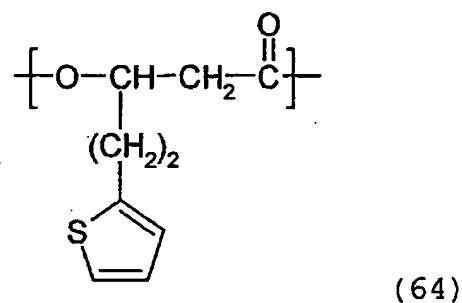
- 95 -

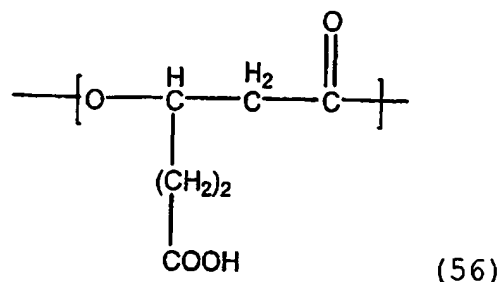
Then, in an iced bath, 420 mg of potassium permanganate were slowly added and an agitation was carried out for 2 hours in an iced bath and 18 hours at the room temperature. After the reaction, 100 ml of water and 1000 mg of sodium bisulfite were added. Then the liquid was brought to pH = 1 by 1.0 N hydrochloric acid. After dichloromethane in the mixed solvent was distilled off in an evaporator, a polymer in the solution was recovered. The polymer was recovered by washing with 200 ml of purified water, then with 200 ml of methanol, three times with 200 ml of purified water and finally with 200 ml of methanol. The obtained polymer was dissolved in 10 ml of tetrahydrofuran and dialyzed for 1 day with a dialysis film (manufactured by Spectrum Inc., Stectra/Por Standard Regenerated Cellulose Dialysis Membrane 3), in a 1-L beaker containing 500 ml of methanol. The polymer present in the dialysis film was recovered and dried under a reduced pressure to obtain 946 mg of a desired PHA.

An average molecular weight of the obtained PHA was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a result there were obtained a number-averaged molecular weight $M_n = 45000$ and a weight-averaged molecular weight $M_w = 95000$.

For specifying the structure of the obtained PHA, a NMR analysis was carried out under conditions same as in Example 1.

As a result, there was confirmed a polyhydroxy
 5 alkananoate copolymer including, as monomer units, 3-hydroxy-5-(2-thienyl)valeric acid represented by the following chemical formula (64), 3-hydroxy-9-carboxynonanoic acid represented by a chemical formula (54), 3-hydroxy-7-carboxyheptanoic acid
 10 represented by a chemical formula (55) and 3-hydroxy-5-carboxyvaleric acid represented by a chemical formula (56).





Also a proportion of the units of the obtained PHA was calculated by a methylesterification, utilizing trimethylsilyldiazomethane, of a carboxyl group at an end of a side chain of the PHA.

30 mg of the object PHA were charged in a 100-ml eggplant-shaped flask and were dissolved by adding 2.1 ml of chloroform and 0.7 ml of methanol. The solution was added with 0.3 ml of a 2.0 mol/L solution of trimethylsilyldiazomethane in hexane (supplied by Aldrich Inc.) and was agitated for 30 minutes at the room temperature. After the reaction, the solvent was distilled off in an evaporator to recover a polymer. The polymer was recovered by washing with 50 ml of methanol. A drying under a reduced pressure provided 30 mg of PHA.

A NMR analysis was carried out as in Example 1. As a result, ¹H-NMR spectrum confirmed a proportion of the units in which 3-hydroxy-5-(2-thienyl)valeric acid was present by 85 mol%, a sum of three units of 3-hydroxy-9-carboxynonanoic acid, 3-hydroxy-7-carboxyheptanoic acid and 3-hydroxy-5-carboxyvaleric acid by 10 mol%, and others (linear 3-hydroxyalkanoic

- 98 -

acid of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or 12 carbon atoms) by 5 mol%.

[Example 18]

There were prepared three 2000-ml shake flasks,
5 and, in each, 0.5 wt.% of polypeptone (supplied by Wako Pure Chemical Co.), 6 mmol/L of 5-(2-thienylsulfanyl)valeric acid, and 1 mmol/L of 10-undecenoic acid were dissolved in 1000 ml of an
aforementioned M9 culture medium, which was placed in
10 a 2000 ml shake flask, then sterilized in an autoclave and cooled to the room temperature. Then 10 ml of a culture liquid of *Pseudomonas cichorii* YN2 strain, shake cultured in advance in an M9 culture medium containing 0.5% of polypeptone for 8 hours,
15 was added to each prepared culture medium, and culture was conducted for 60 hours at 30°C. After the culture, the culture liquids were united, and the cells were recovered by centrifuging, rinsed with methanol and dried. The dried cells, after weighing,
20 were agitated with chloroform for 72 hours at 25°C to extract a polymer. The chloroform extract was filtered with a 0.45 µm membrane filter, then concentrated in an evaporator, and the polymer was recovered by a reprecipitation in cold methanol. A
25 desired polymer was then obtained by drying under a reduced pressure.

According to a weighing of the obtained polymer,

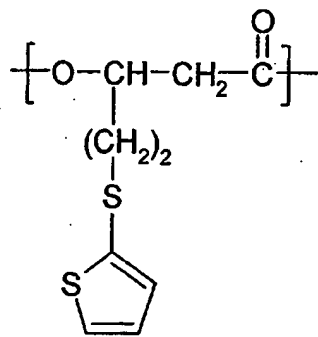
- 99 -

1257 mg (dry weight) of PHA were obtained in the present example.

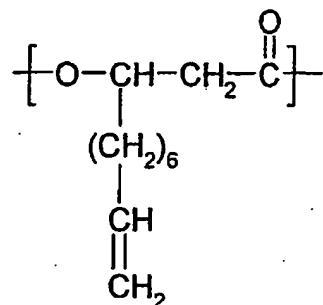
An average molecular weight of the obtained PHA was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a result there were obtained a number-averaged molecular weight $M_n = 68000$ and a weight-averaged molecular weight $M_w = 160000$.

For specifying the structure of the obtained PHA, a NMR analysis was conducted under conditions same as in Example 1.

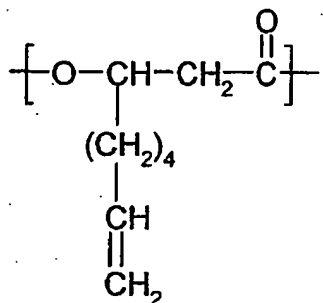
As a result, there was confirmed a polyhydroxy alkanoate copolymer including, as monomer units, 3-hydroxy-5-(2-thienylsulfanyl)valeric acid represented by the following chemical formula (65), 3-hydroxy-10-undecenoic acid represented by a chemical formula (5), 3-hydroxy-8-nonenoic acid represented by a chemical formula (6) and 3-hydroxy-6-heptenoic acid represented by a chemical formula (7).



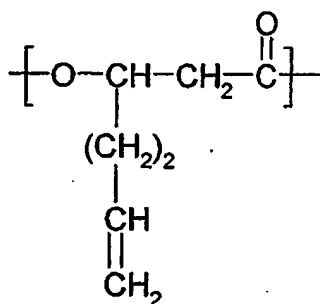
(65)



(5)



(6)



(7)

Also a proportion of such units was confirmed
 5 by ^1H -NMR spectrum, where a sum of three units of 3-
 hydroxy-10-undecenoic acid, 3-hydroxy-8-nonenic acid
 and 3-hydroxy-6-heptenoic acid was present by 9 mol%,
 3-hydroxy-5-(2-thienylsulfanyl)valeric acid by 84
 mol% and others (linear 3-hydroxyalkanoic acid of 4
 10 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with
 10 or 12 carbon atoms) by 7 mol%.

[Example 19]

- 101 -

There were prepared ten 2000-ml shake flasks, and, in each, 0.5 wt.% of polypeptone (supplied by Wako Pure Chemical Co.), 6 mmol/L of 5-(2-thienylcarbonyl)valeric acid, and 1 mmol/L of 10-undecenoic acid were dissolved in 1000 ml of an aforementioned M9 culture medium, which was placed in a 2000 ml shake flask, then sterilized in an autoclave and cooled to the room temperature. Then 10 ml of a culture liquid of *Pseudomonas cichorii* YN2 strain, shake cultured in advance in an M9 culture medium containing 0.5% of polypeptone for 8 hours, was added to each prepared culture medium, and culture was conducted for 60 hours at 30°C. After the culture, the culture liquids were united, and the cells were recovered by centrifuging, rinsed with methanol and dried. The dried cells, after weighing, were agitated with chloroform for 72 hours at 25°C to extract a polymer. The chloroform extract was filtered with a 0.45 μ m membrane filter, then concentrated in an evaporator, and the polymer was recovered by a reprecipitation in cold methanol. A desired polymer was then obtained by drying under a reduced pressure.

According to a weighing of the obtained polymer, 1251 mg (dry weight) of PHA were obtained in the present example.

An average molecular weight of the obtained PHA

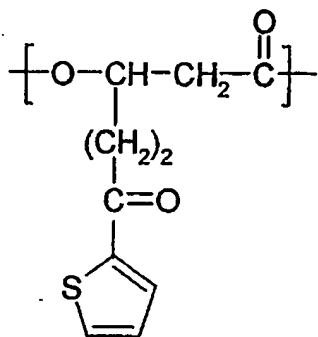
was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a result there were obtained a number-averaged

5 molecular weight $M_n = 75000$ and a weight-averaged molecular weight $M_w = 180000$.

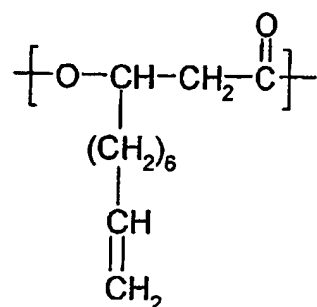
For specifying the structure of the obtained PHA, a NMR analysis was conducted under conditions similar to those in Example 1.

10 As a result, there was confirmed a polyhydroxy alkanate copolymer including, as monomer units, 3-hydroxy-5-(2-thienylcarbonyl)valeric acid represented by the following chemical formula (66), 3-hydroxy-10-undecenoic acid represented by a chemical formula (5),

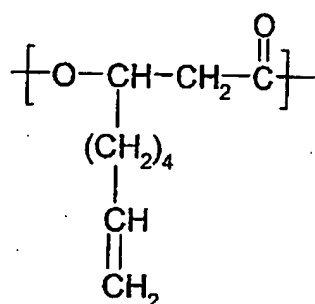
15 3-hydroxy-8-nonenoic acid represented by a chemical formula (6) and 3-hydroxy-6-heptenoic acid represented by a chemical formula (7).



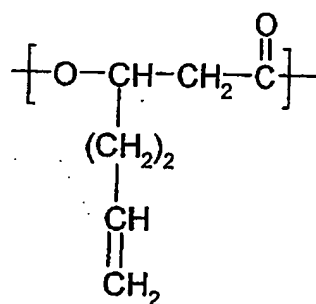
(66)



(5)



(6)



(7)

Also a proportion of such units was confirmed
 5 by ^1H -NMR spectrum, where a sum of three units of 3-
 hydroxy-10-undecenoic acid, 3-hydroxy-8-nonenoic acid
 and 3-hydroxy-6-heptenoic acid was present by 10 mol%,
 3-hydroxy-5-(2-thienylcarbonyl)valeric acid by 81
 mol% and others (linear 3-hydroxyalkanoic acid of 4
 10 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with
 10 or 12 carbon atoms) by 9 mol%.

The polyhydroxy alkanoate thus obtained was

- 104 -

utilized in the following reaction.

999 mg of polyhydroxy alkanate were charged in a 500-ml eggplant-shaped flask and were dissolved by adding 60 ml of dichloromethane. The solution was
5 placed in an iced bath, and 10 ml of acetic acid and 382 mg of 18-crown-6-ether were added and agitated. Then, in an iced bath, 304 mg of potassium permanganate were slowly added and an agitation was carried out for 2 hours in an iced bath and 18 hours
10 at the room temperature. After the reaction, 100 ml of water and 1000 mg of sodium bisulfite were added. Then the liquid was brought to pH = 1 by 1.0 N hydrochloric acid. After dichloromethane in the mixed solvent was distilled off in an evaporator, a
15 polymer in the solution was recovered. The polymer was recovered by washing with 200 ml of purified water, then with 200 ml of methanol, three times with 200 ml of purified water and finally with 200 ml of methanol. The obtained polymer was dissolved in 10
20 ml of tetrahydrofuran and dialyzed for 1 day with a dialysis film (manufactured by Spectrum Inc., Stectra/Por Standard Regenerated Cellulose Dialysis Membrane 3), in a 1-L beaker containing 500 ml of methanol. The polymer present in the dialysis film
25 was recovered and dried under a reduced pressure to obtain 935 mg of a desired PHA.

An average molecular weight of the obtained PHA

- 105 -

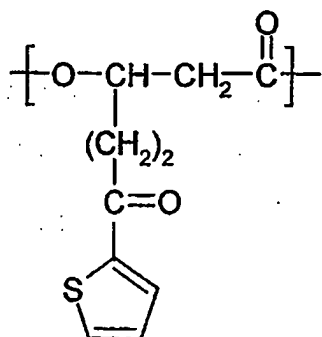
was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a result there were obtained a number-averaged

5 molecular weight $M_n = 45000$ and a weight-averaged molecular weight $M_w = 99000$.

For specifying the structure of the obtained PHA, a NMR analysis was carried out under conditions same as in Example 1.

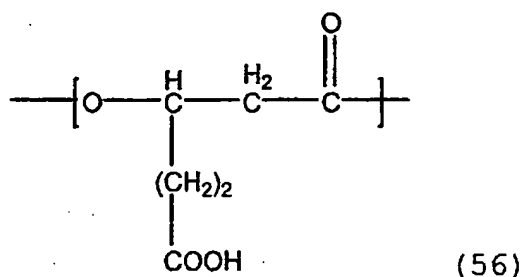
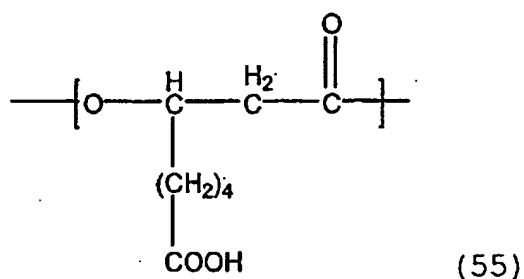
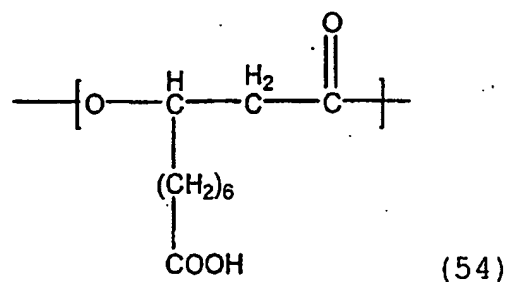
10 As a result, there was confirmed a polyhydroxy alkanoate copolymer including, as monomer units, 3-hydroxy-5-(2-thienylcarbonyl)valeric acid represented by the following chemical formula (66), 3-hydroxy-9-carboxynonanoic acid represented by a chemical

15 formula (54), 3-hydroxy-7-carboxyheptanoic acid represented by a chemical formula (55) and 3-hydroxy-5-carboxyvaleric acid represented by a chemical formula (56).



(66)

- 106 -



Also a proportion of the units of the obtained
 5 PHA was calculated by a methylesterification,
 utilizing trimethylsilyldiazomethane, of a carboxyl
 group at an end of a side chain of the PHA.

31 mg of the object PHA were charged in a 100-
 ml eggplant-shaped flask and were dissolved by adding
 10 2.1 ml of chloroform and 0.7 ml of methanol. The
 solution was added with 0.3 ml of a 2.0 mol/L
 solution of trimethylsilyldiazomethane in hexane
 (supplied by Aldrich Inc.) and was agitated for 30
 minutes at the room temperature. After the reaction,
 15 the solvent was distilled off in an evaporator to

- 107 -

recover a polymer. The polymer was recovered by washing with 50 ml of methanol. A drying under a reduced pressure provided 30 mg of PHA.

A NMR analysis was carried out as in Example 1.

- 5 As a result, ^1H -NMR spectrum confirmed a proportion of the units in which 3-hydroxy-5-(2-thienylcarbonyl)valeric acid was present by 83 mol%, a sum of three units of 3-hydroxy-9-carboxynonanoic acid, 3-hydroxy-7-carboxyheptanoic acid and 3-
10 hydroxy-5-carboxyvaleric acid by 7 mol%, and others (linear 3-hydroxyalkanoic acid of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or 12 carbon atoms) by 10 mol%.

[Example 20]

- 15 There were prepared fifteen 2000-ml shake flasks, and, in each, 0.5 wt.% of polypeptone (supplied by Wako Pure Chemical Co.), 6 mmol/L of 5-[(phenylmethyl)oxy] valeric acid, and 1 mmol/L of 10-undecenoic acid were dissolved in 1000 ml of an
20 aforementioned M9 culture medium, which was placed in a 2000 ml shake flask, then sterilized in an autoclave and cooled to the room temperature. Then 10 ml of a culture liquid of *Pseudomonas cichorii* YN2 strain, shake cultured in advance in an M9 culture
25 medium containing 0.5% of polypeptone for 8 hours, was added to each prepared culture medium, and culture was conducted for 60 hours at 30°C. After

- 108 -

the culture, the culture liquids were united, and the cells were recovered by centrifuging, rinsed with methanol and dried. The dried cells, after weighing, were agitated with chloroform for 72 hours at 25°C to
5 extract a polymer. The chloroform extract was filtered with a 0.45 μ m membrane filter, then concentrated in an evaporator, and the polymer was recovered by a reprecipitation in cold methanol. A desired polymer was then obtained by drying under a
10 reduced pressure.

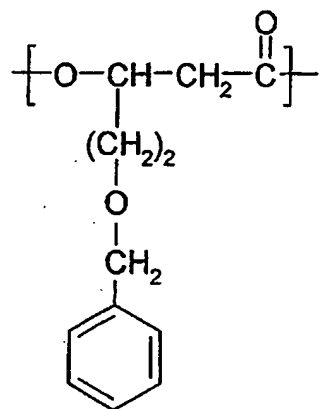
According to a weighing of the obtained polymer, 1348 mg (dry weight) of PHA were obtained in the present example.

An average molecular weight of the obtained PHA
15 was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a result there were obtained a number-averaged molecular weight $M_n = 79000$ and a weight-averaged
20 molecular weight $M_w = 190000$.

For specifying the structure of the obtained PHA, a NMR analysis was conducted under conditions similar to those in Example 1.

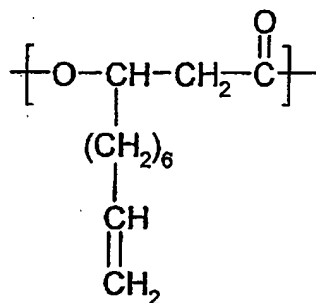
As a result, there was confirmed a polyhydroxy
25 alkanoate copolymer including, as monomer units, 3-hydroxy-5-[(phenylmethyl)oxy]valeric acid represented by the following chemical formula (67), 3-hydroxy-10-

undecenoic acid represented by a chemical formula (5),
3-hydroxy-8-nonenoic acid represented by a chemical
formula (6) and 3-hydroxy-6-heptenoic acid
represented by a chemical formula (7).

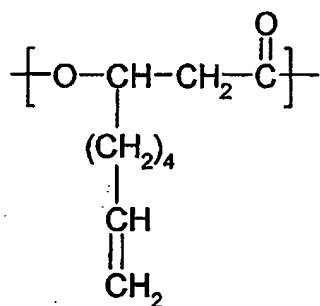


5

(67)

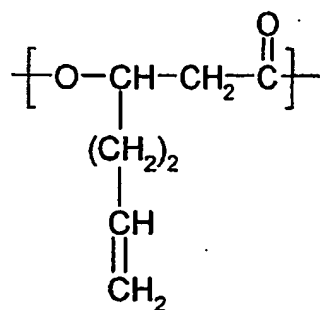


(5)



(6)

- 110 -



(7)

Also a proportion of such units was confirmed by ^1H -NMR spectrum, where a sum of three units of 3-hydroxy-10-undecenoic acid, 3-hydroxy-8-nonenoic acid and 3-hydroxy-6-heptenoic acid was present by 10 mol%, 3-hydroxy-5-[(phenylmethyl)oxy]valeric acid by 82 mol% and others (linear 3-hydroxyalkanoic acid of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or 12 carbon atoms) by 8 mol%.

The polyhydroxy alkanoate thus obtained was utilized in the following reaction.

1004 mg of polyhydroxy alkanoate were charged in a 500-ml eggplant-shaped flask and were dissolved by adding 60 ml of dichloromethane. The solution was placed in an iced bath, and 10 ml of acetic acid and 389 mg of 18-crown-6-ether were added and agitated. Then, in an iced bath, 310 mg of potassium permanganate were slowly added and an agitation was carried out for 2 hours in an iced bath and 18 hours at the room temperature. After the reaction, 100 ml of water and 1000 mg of sodium bisulfite were added. Then the liquid was brought to pH = 1 by 1.0 N

- 111 -

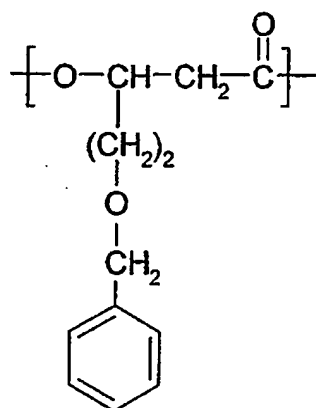
hydrochloric acid. After dichloromethane in the mixed solvent was distilled off in an evaporator, a polymer in the solution was recovered. The polymer was recovered by washing with 200 ml of purified
5 water, then with 200 ml of methanol, three times with 200 ml of purified water and finally with 200 ml of methanol. The obtained polymer was dissolved in 10 ml of tetrahydrofuran and dialyzed for 1 day with a dialysis film (manufactured by Spectrum Inc.,
10 Stectra/Por Standard Regenerated Cellulose Dialysis Membrane 3), in a 1-L beaker containing 500 ml of methanol. The polymer present in the dialysis film was recovered and dried under a reduced pressure to obtain 940 mg of a desired PHA.

15 An average molecular weight of the obtained PHA was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a result there were obtained a number-averaged
20 molecular weight $M_n = 48000$ and a weight-averaged molecular weight $M_w = 106000$.

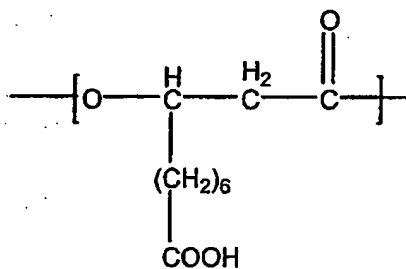
For specifying the structure of the obtained PHA, a NMR analysis was carried out under conditions same as in Example 1.

25 As a result, there was confirmed a polyhydroxy alkanoate copolymer including, as monomer units, 3-hydroxy-5-[(phenylmethyl)oxy]valeric acid represented

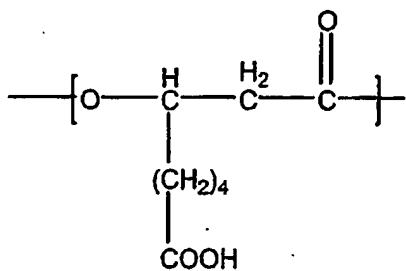
by the following chemical formula (67), 3-hydroxy-9-carboxynonanoic acid represented by a chemical formula (54), 3-hydroxy-7-carboxyheptanoic acid represented by a chemical formula (55) and 3-hydroxy-5-carboxyvaleric acid represented by a chemical formula (56).



(67)

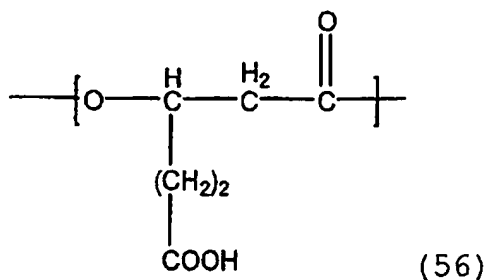


(54)



(55)

- 113 -



Also a proportion of the units of the obtained PHA was calculated by a methylesterification, utilizing trimethylsilyldiazomethane, of a carboxyl group at an end of a side chain of the PHA.

30 mg of the object PHA were charged in a 100-ml eggplant-shaped flask and were dissolved by adding 2.1 ml of chloroform and 0.7 ml of methanol. The solution was added with 0.3 ml of a 2.0 mol/L solution of trimethylsilyldiazomethane in hexane (supplied by Aldrich Inc.) and was agitated for 30 minutes at the room temperature. After the reaction, the solvent was distilled off in an evaporator to recover a polymer. The polymer was recovered by washing with 50 ml of methanol. A drying under a reduced pressure provided 29 mg of PHA.

A NMR analysis was carried out as in Example 1. As a result, ¹H-NMR spectrum confirmed a proportion of the units in which 3-hydroxy-5-[(phenylmethyl)oxy] valeric acid was present by 84 mol%, a sum of three units of 3-hydroxy-9-carboxynonanoic acid, 3-hydroxy-7-carboxyheptanoic acid and 3-hydroxy-5-carboxyvaleric acid by 8 mol%, and others (linear 3-

- 114 -

hydroxyalkanoic acid of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or 12 carbon atoms) by 8 mol%.

[Example 21]

5 There were prepared five 2000-ml shake flasks, and, in each, 0.5 wt.% of polypeptone (supplied by Wako Pure Chemical Co.), 3 mmol/L of 5-phenoxyvaleric acid, 3 mmol/L of 5-cyclohexylvaleric acid and 1 mmol/L of 10-undecenoic acid were dissolved in 1000
10 ml of an aforementioned M9 culture medium, which was placed in a 2000 ml shake flask, then sterilized in an autoclave and cooled to the room temperature. Then 10 ml of a culture liquid of *Pseudomonas cichorii* YN2 strain, shake cultured in advance in an
15 M9 culture medium containing 0.5% of polypeptone for 8 hours, was added to each prepared culture medium, and culture was conducted for 60 hours at 30°C. After the culture, the culture liquids were united, and the cells were recovered by centrifuging, rinsed
20 with methanol and dried. The dried cells, after weighing, were agitated with chloroform for 72 hours at 25°C to extract a polymer. The chloroform extract was filtered with a 0.45 μ m membrane filter, then concentrated in an evaporator, and the polymer was
25 recovered by a reprecipitation in cold methanol. A desired polymer was then obtained by drying under a reduced pressure.

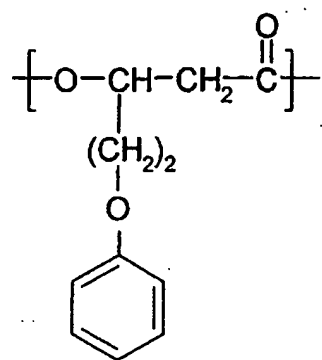
- 115 -

According to a weighing of the obtained polymer, 1285 mg (dry weight) of PHA were obtained in the present example.

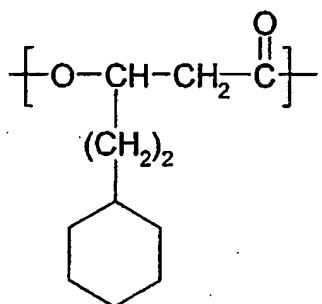
An average molecular weight of the obtained PHA was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a result there were obtained a number-averaged molecular weight $M_n = 86000$ and a weight-averaged molecular weight $M_w = 230000$.

For specifying the structure of the obtained PHA, a NMR analysis was conducted under conditions similar to those in Example 1.

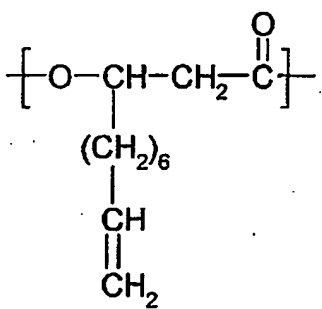
As a result, there was confirmed a polyhydroxy alkanooate copolymer including, as monomer units, 3-hydroxy-5-phenoxyvaleric acid represented by the following chemical formula (53), 3-hydroxy-5-cyclohexylvaleric acid represented by the following chemical formula (68), 3-hydroxy-10-undecenoic acid represented by a chemical formula (5), 3-hydroxy-8-nonenic acid represented by a chemical formula (6) and 3-hydroxy-6-heptenoic acid represented by a chemical formula (7).



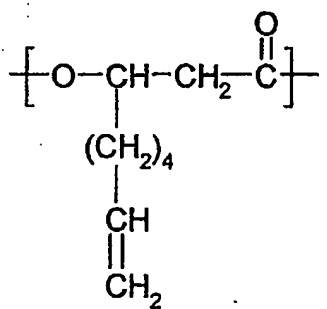
(53)



(68)

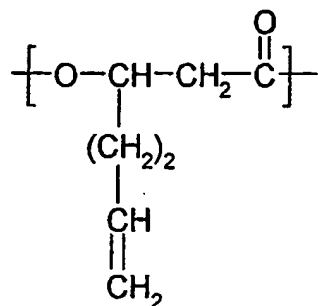


(5)



(6)

- 117 -



(7)

Also a proportion of such units was confirmed by ^1H -NMR spectrum, where a sum of three units of 3-hydroxy-10-undecenoic acid, 3-hydroxy-8-nonenoic acid and 3-hydroxy-6-heptenoic acid was present by 7 mol%,
 5 3-hydroxy-5-phenoxyvaleric acid by 48 mol%, 3-hydroxy-5-cyclohexylvaleric acid by 41 mol%,
 and others (linear 3-hydroxyalkanoic acid of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or
 10 12 carbon atoms) by 4 mol%.

The polyhydroxy alkanoate thus obtained was utilized in the following reaction.

1002 mg of polyhydroxy alkanoate were charged in a 500-ml eggplant-shaped flask and were dissolved
 15 by adding 60 ml of dichloromethane. The solution was placed in an iced bath, and 10 ml of acetic acid and 288 mg of 18-crown-6-ether were added and agitated.
 Then, in an iced bath, 230 mg of potassium permanganate were slowly added and an agitation was
 20 carried out for 2 hours in an iced bath and 18 hours at the room temperature. After the reaction, 100 ml of water and 1000 mg of sodium bisulfite were added.

- 118 -

Then the liquid was brought to pH = 1 by 1.0 N hydrochloric acid. After dichloromethane in the mixed solvent was distilled off in an evaporator, a polymer in the solution was recovered. The polymer
5 was recovered by washing with 200 ml of purified water, then with 200 ml of methanol, three times with 200 ml of purified water and finally with 200 ml of methanol. The obtained polymer was dissolved in 10 ml of tetrahydrofuran and dialyzed for 1 day with a
10 dialysis film (manufactured by Spectrum Inc., Stectra/Por Standard Regenerated Cellulose Dialysis Membrane 3), in a 1-L beaker containing 500 ml of methanol. The polymer present in the dialysis film was recovered and dried under a reduced pressure to
15 obtain 967 mg of a desired PHA.

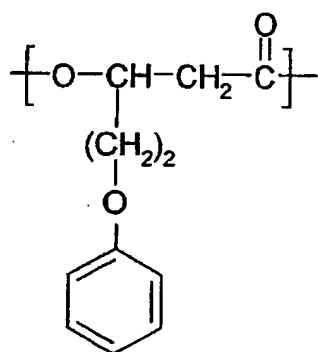
An average molecular weight of the obtained PHA was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a
20 result there were obtained a number-averaged molecular weight $M_n = 51000$ and a weight-averaged molecular weight $M_w = 108000$.

For specifying the structure of the obtained PHA, a NMR analysis was carried out under conditions
25 same as in Example 1.

As a result, there was confirmed a polyhydroxy alkanoate copolymer including, as monomer units, 3-

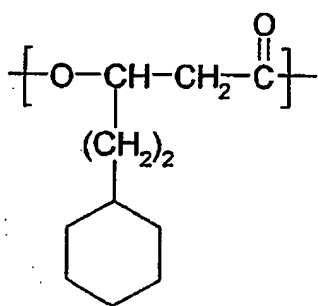
- 119 -

hydroxy-5-phenoxyvaleric acid represented by the following chemical formula (53), 3-hydroxy-5-cyclohexylvaleric acid represented by the following chemical formula (68), 3-hydroxy-9-carboxynonanoic acid represented by a chemical formula (54), 3-hydroxy-7-carboxyheptanoic acid represented by a chemical formula (55) and 3-hydroxy-5-carboxyvaleric acid represented by a chemical formula (56).

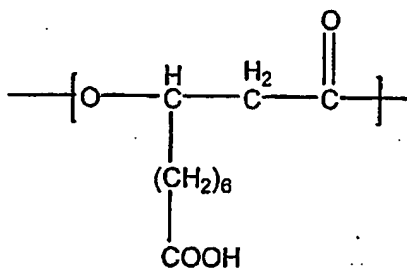


10

(53)

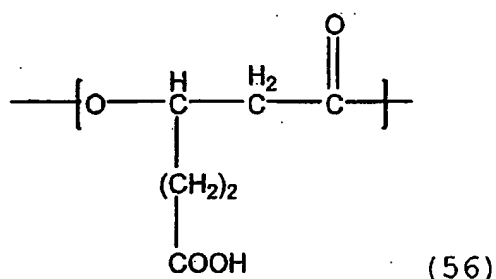
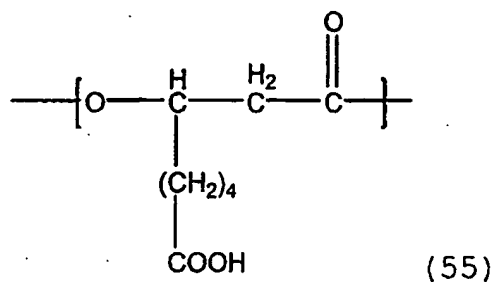


(68)



(54)

- 120 -



Also a proportion of the units of the obtained PHA was calculated by a methylesterification, utilizing trimethylsilyldiazomethane, of a carboxyl group at an end of a side chain of the PHA.

30 mg of the object PHA were charged in a 100-ml eggplant-shaped flask and were dissolved by adding 2.1 ml of chloroform and 0.7 ml of methanol. The solution was added with 0.3 ml of a 2.0 mol/L solution of trimethylsilyldiazomethane in hexane (supplied by Aldrich Inc.) and was agitated for 30 minutes at the room temperature. After the reaction, the solvent was distilled off in an evaporator to recover a polymer. The polymer was recovered by washing with 50 ml of methanol. A drying under a reduced pressure provided 28 mg of PHA.

A NMR analysis was carried out as in Example 1. As a result, ^1H -NMR spectrum confirmed a proportion

- 121 -

of the units in which 3-hydroxy-5-phenoxyvaleric acid was present by 49 mol%, 3-hydroxy-5-cyclohexylvaleric acid by 42 mol%, a sum of three units of 3-hydroxy-9-carboxynonanoic acid, 3-hydroxy-7-carboxyheptanoic acid and 3-hydroxy-5-carboxyvaleric acid by 6 mol%, and others (linear 3-hydroxyalkanoic acid of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or 12 carbon atoms) by 3 mol%.

[Example 22]

10 There were prepared two 2000-ml shake flasks, and, in each, 0.5 wt.% of polypeptone (supplied by Wako Pure Chemical Co.), 4 mmol/L of 5-phenylvaleric acid, and 1 mmol/L of dodecanedioic acid monoethyl ester were dissolved in 1000 ml of an aforementioned
15 M9 culture medium, which was placed in a 2000 ml shake flask, then sterilized in an autoclave and cooled to the room temperature. Then 5 ml of a culture liquid of *Pseudomonas cichorii* YN2 strain, shake cultured in advance in an M9 culture medium
20 containing 0.5% of polypeptone for 8 hours at 30°C, was added to each prepared culture medium, and culture was conducted for 41 hours at 30°C. After the culture, the cells were recovered by centrifuging, rinsed with methanol and lyophilized. The dried
25 cells, after weighing, were agitated with chloroform for 48 hours at 50°C to extract a polymer. The chloroform extract was filtered, then concentrated in

- 122 -

an evaporator, and a solid precipitate formed with cold methanol was collected and dried under a reduced pressure to obtain a desired polymer. According to a weighing of the obtained polymer, 910 mg (dry weight) of PHA were obtained in the present example.

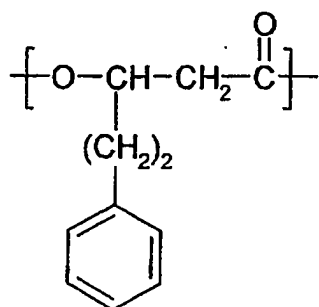
An average molecular weight of the obtained PHA was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a result there were obtained a number-averaged molecular weight $M_n = 78000$ and a weight-averaged molecular weight $M_w = 157000$.

For specifying the structure of the obtained PHA, a NMR analysis was conducted under same conditions as in Example 1.

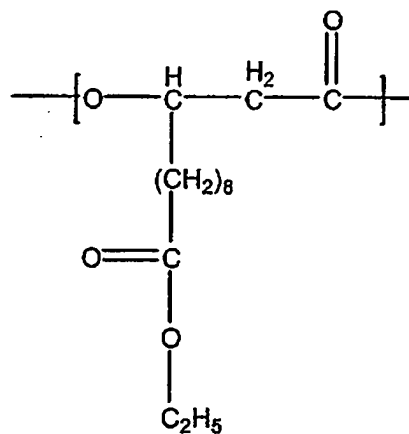
As a result, there was confirmed a polyhydroxy alkanooate copolymer including, as monomer units, 3-hydroxy-5-phenylvaleric acid represented by the following chemical formula (60) by 78 mol%, three units of 3-hydroxy-11-ethoxycarbonylundecanoic acid represented by the following chemical formula (69), 3-hydroxy-9-ethoxycarbonylnonanoic acid represented by a chemical formula (70), and 3-hydroxy-7-ethoxycarbonylheptanoic acid represented by a chemical formula (71) collectively by 14 mol%, and others (linear 3-hydroxyalkanoic acid of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or

- 123 -

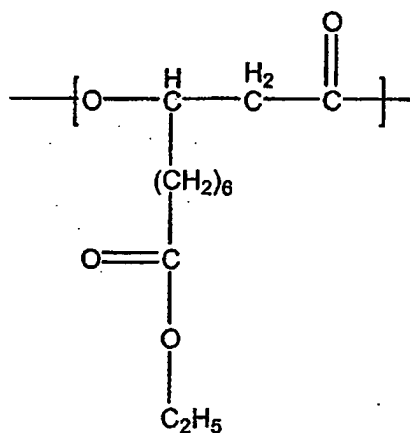
12 carbon atoms) by 8 mol%.



(60)

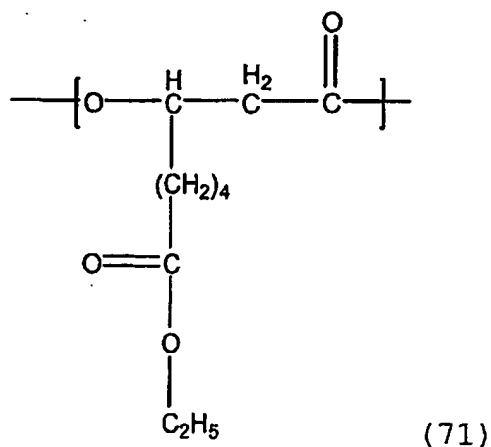


(69)



(70)

- 124 -



[Example 23]

There were prepared two 2000-ml shake flasks, and, in each, 0.5 wt.% of yeast extract (supplied by

5 DIFCO), 4 mmol/L of 5-phenylvaleric acid, and 1 mmol/L of dodecanedioic acid monoethyl ester were dissolved in 1000 ml of an aforementioned M9 culture medium, which was placed in a 2000 ml shake flask, then sterilized in an autoclave and cooled to the

10 room temperature. Then 5 ml of a culture liquid of *Pseudomonas cichorii* YN2 strain, shake cultured in advance in an M9 culture medium containing 0.5% of polypeptone for 8 hours at 30°C, was added to each prepared culture medium, and culture was conducted

15 for 40 hours at 30°C. After the culture, the cells were recovered by centrifuging, rinsed with methanol and lyophilized. The dried cells, after weighing, were agitated with chloroform for 48 hours at 50°C to extract a polymer. The chloroform extract was

20 filtered, then concentrated in an evaporator, and a

- 125 -

solid precipitate formed with cold methanol was collected and dried under a reduced pressure to obtain a desired polymer. According to a weighing of the obtained polymer, 250 mg (dry weight) of PHA were
5 obtained in the present example.

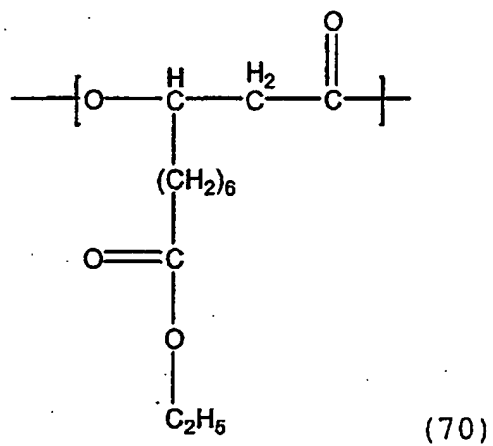
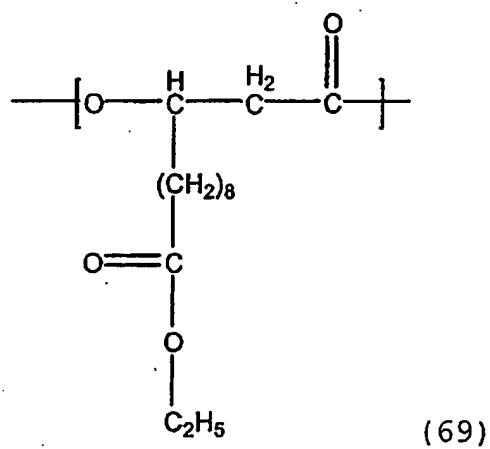
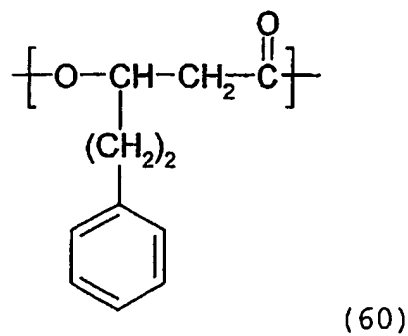
An average molecular weight of the obtained PHA was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a
10 result there were obtained a number-averaged molecular weight $M_n = 75000$ and a weight-averaged molecular weight $M_w = 152000$.

For specifying the structure of the obtained PHA, a NMR analysis was conducted under same
15 conditions as in Example 1.

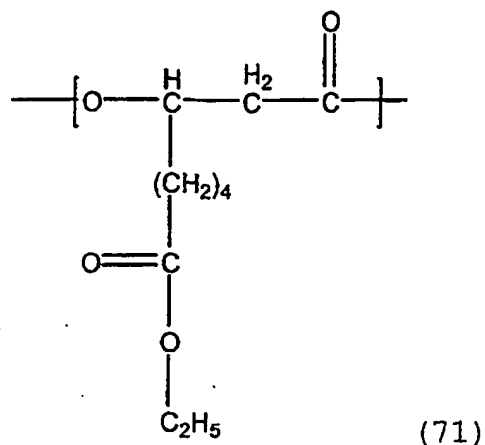
As a result, there was confirmed a polyhydroxy alkanate copolymer including, as monomer units, 3-hydroxy-5-phenylvaleric acid represented by the following chemical formula (60) by 75 mol%, three
20 units of 3-hydroxy-11-ethoxycarbonylundecanoic acid represented by the following chemical formula (69), 3-hydroxy-9-ethoxycarbonylnonanoic acid represented by a chemical formula (70), and 3-hydroxy-7-ethoxycarbonylheptanoic acid represented by a
25 chemical formula (71) collectively by 15 mol%, and others (linear 3-hydroxyalkanoic acid of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or

- 126 -

12 carbon atoms) by 10 mol%.



- 127 -



[Example 24]

There were prepared two 2000-ml shake flasks, and, in each, 0.5 wt.% of D-glucose (supplied by Kishida Kagaku), 4 mmol/L of 5-phenylvaleric acid, and 1 mmol/L of dodecanedioic acid monoethyl ester were dissolved in 1000 ml of an aforementioned M9 culture medium, which was placed in a 2000 ml shake flask, then sterilized in an autoclave and cooled to the room temperature. Then 5 ml of a culture liquid of *Pseudomonas jessenii* P161 strain, shake cultured in advance in an M9 culture medium containing 0.5% of polypeptone for 8 hours at 30°C, was added to each prepared culture medium, and culture was conducted for 40 hours at 30°C. After the culture, the cells were recovered by centrifuging, rinsed with methanol and lyophilized. The dried cells, after weighing, were agitated with chloroform for 48 hours at 50°C to extract a polymer. The chloroform extract was filtered, then concentrated in an evaporator, and a

- 128 -

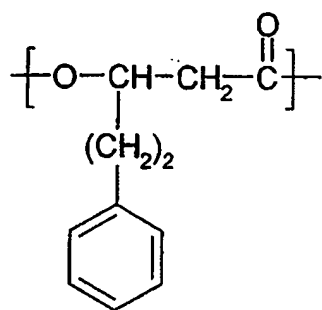
solid precipitate formed with cold methanol was collected and dried under a reduced pressure to obtain a desired polymer. According to a weighing of the obtained polymer, 300 mg (dry weight) of PHA were
5 obtained in the present example.

An average molecular weight of the obtained PHA was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a
10 result there were obtained a number-averaged molecular weight $M_n = 71000$ and a weight-averaged molecular weight $M_w = 149000$.

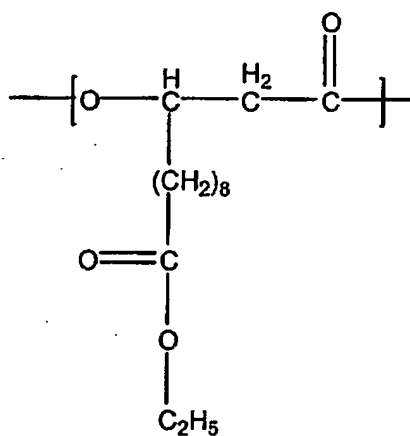
For specifying the structure of the obtained PHA, a NMR analysis was conducted under same
15 conditions as in Example 1.

As a result, there was confirmed a polyhydroxy alkanate copolymer including, as monomer units, 3-hydroxy-5-phenylvaleric acid represented by the following chemical formula (60) by 78 mol%, three
20 units of 3-hydroxy-11-ethoxycarbonylundecanoic acid represented by the following chemical formula (69), 3-hydroxy-9-ethoxycarbonylnonanoic acid represented by a chemical formula (70), and 3-hydroxy-7-ethoxycarbonylheptanoic acid represented by a
25 chemical formula (71) collectively by 14 mol%, and others (linear 3-hydroxyalkanoic acid of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or

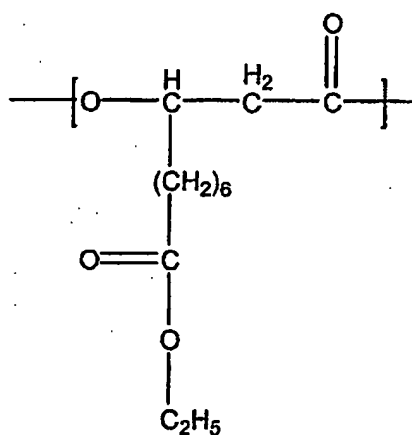
12 carbon atoms) by 8 mol%.



(60)

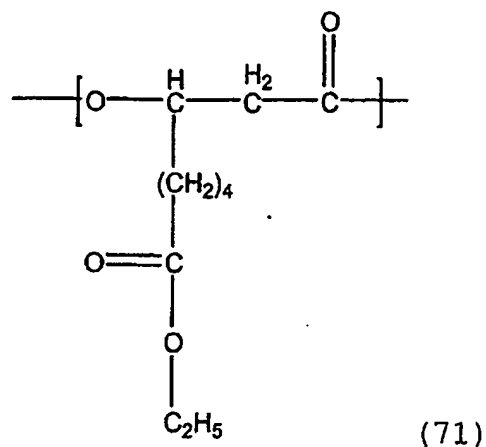


(69)



(70)

- 130 -



[Example 25]

There were prepared two 2000-ml shake flasks, and, in each, 0.5 wt.% of polypeptone (supplied by Wako Pure Chemicals Co.), 4 mmol/L of 5-phenoxyvaleric acid, and 1 mmol/L of dodecanedioic acid monoethyl ester were dissolved in 1000 ml of an aforementioned M9 culture medium, which was placed in a 2000 ml shake flask, then sterilized in an autoclave and cooled to the room temperature. Then 5 ml of a culture liquid of *Pseudomonas cichorii* YN2 strain, shake cultured in advance in an M9 culture medium containing 0.5% of polypeptone for 8 hours at 30°C, was added to each prepared culture medium, and culture was conducted for 41 hours at 30°C. After the culture, the cells were recovered by centrifuging, rinsed with methanol and lyophilized. The dried cells, after weighing, were agitated with chloroform for 48 hours at 50°C to extract a polymer. The chloroform extract was filtered, then concentrated in

- 131 -

an evaporator, and a solid precipitate formed with cold methanol was collected and dried under a reduced pressure to obtain a desired polymer. According to a weighing of the obtained polymer, 680 mg (dry weight) of PHA were obtained in the present example.

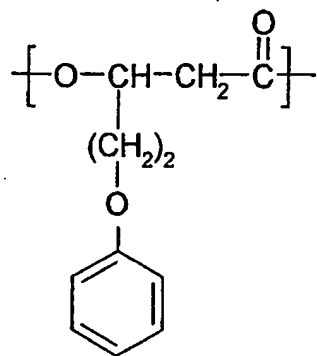
An average molecular weight of the obtained PHA was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a result there were obtained a number-averaged molecular weight $M_n = 69000$ and a weight-averaged molecular weight $M_w = 135000$.

For specifying the structure of the obtained PHA, a NMR analysis was conducted under same conditions as in Example 1.

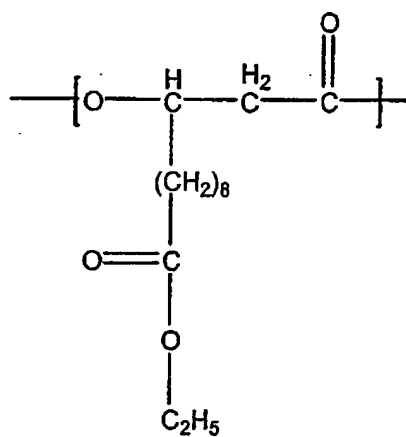
As a result, there was confirmed a polyhydroxyalkanoate copolymer including, as monomer units, 3-hydroxy-5-phenoxyvaleric acid represented by the following chemical formula (53) by 74 mol%, three units of 3-hydroxy-11-ethoxycarbonylundecanoic acid represented by the following chemical formula (69), 3-hydroxy-9-ethoxycarbonylnonanoic acid represented by a chemical formula (70), and 3-hydroxy-7-ethoxycarbonylheptanoic acid represented by a chemical formula (71) collectively by 17 mol%, and others (linear 3-hydroxyalkanoic acid of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or

- 132 -

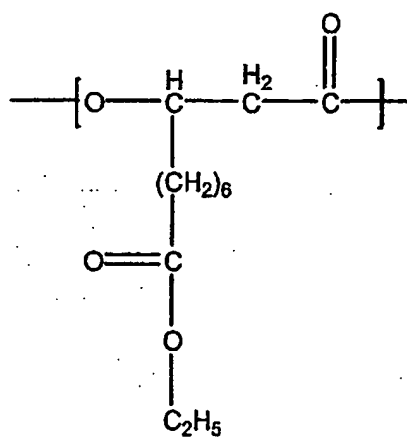
12 carbon atoms) by 9 mol%.



(53)

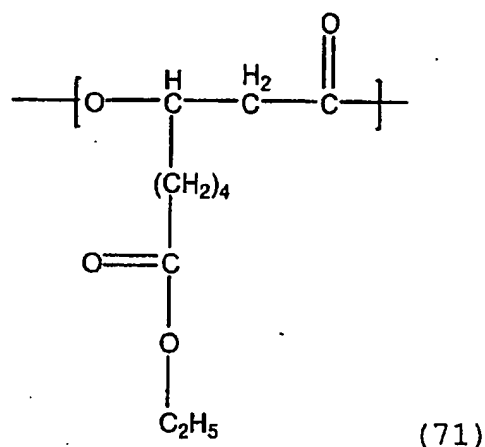


(69)



(70)

- 133 -



[Example 26]

There were prepared two 2000-ml shake flasks, and, in each, 0.5 wt.% of polypeptone (supplied by Wako Pure Chemicals Co.), 4 mmol/L of 4-cyclohexylbutyric acid, and 1 mmol/L of dodecanedioic acid monoethyl ester were dissolved in 1000 ml of an aforementioned M9 culture medium, which was placed in a 2000 ml shake flask, then sterilized in an autoclave and cooled to the room temperature. Then 5 ml of a culture liquid of *Pseudomonas cichorii* YN2 strain, shake cultured in advance in an M9 culture medium containing 0.5% of polypeptone for 8 hours at 30°C, was added to each prepared culture medium, and culture was conducted for 41 hours at 30°C. After the culture, the cells were recovered by centrifuging, rinsed with methanol and lyophilized. The dried cells, after weighing, were agitated with chloroform for 48 hours at 50°C to extract a polymer. The chloroform extract was filtered, then concentrated in

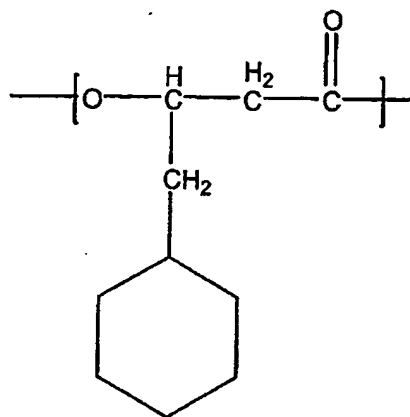
an evaporator, and a solid precipitate formed with cold methanol was collected and dried under a reduced pressure to obtain a desired polymer. According to a weighing of the obtained polymer, 720 mg (dry weight) of PHA were obtained in the present example.

An average molecular weight of the obtained PHA was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a result there were obtained a number-averaged molecular weight $M_n = 81000$ and a weight-averaged molecular weight $M_w = 160000$.

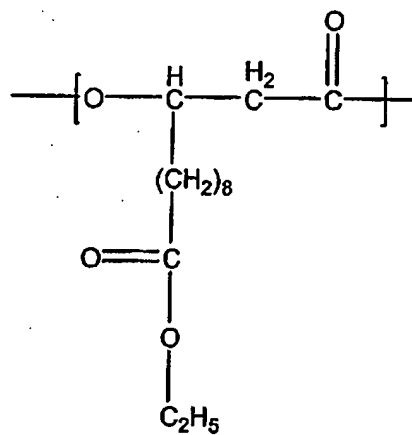
For specifying the structure of the obtained PHA, a NMR analysis was conducted under same conditions as in Example 1.

As a result, there was confirmed a polyhydroxy alkanoate copolymer including, as monomer units, 3-hydroxy-4-cyclohexylbutyric acid represented by the following chemical formula (57) by 76 mol%, three units of 3-hydroxy-11-ethoxycarbonylundecanoic acid represented by the following chemical formula (69), 3-hydroxy-9-ethoxycarbonylnonanoic acid represented by a chemical formula (70), and 3-hydroxy-7-ethoxycarbonylheptanoic acid represented by a chemical formula (71) collectively by 16 mol%, and others (linear 3-hydroxyalkanoic acid of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or

12 carbon atoms) by 8 mol%.

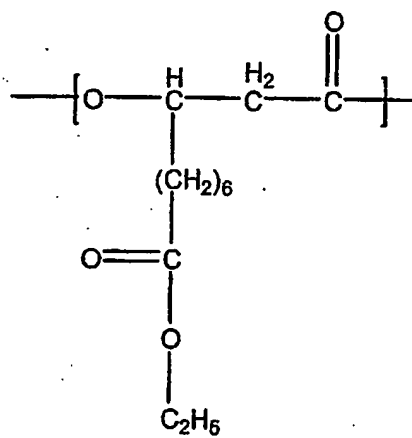


(57)



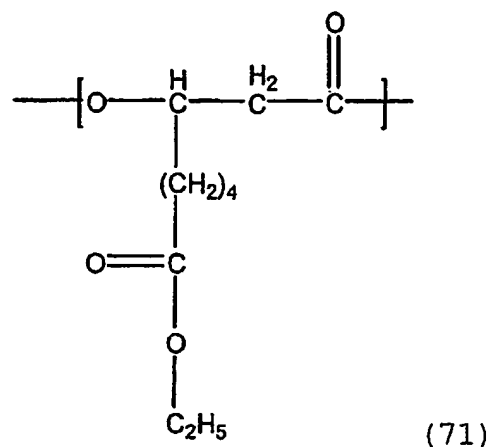
5

(69)



(70)

- 136 -



[Example 27]

There were prepared two 2000-ml shake flasks, and, in each, 0.5 wt.% of polypeptone (supplied by Wako Pure Chemicals Co.), 4 mmol/L of 5-(phenylsulfanyl)valeric acid, and 1 mmol/L of dodecanedioic acid monoethyl ester were dissolved in 1000 ml of an aforementioned M9 culture medium, which was placed in a 2000 ml shake flask, then sterilized in an autoclave and cooled to the room temperature. Then 5 ml of a culture liquid of *Pseudomonas cichorii* YN2 strain, shake cultured in advance in an M9 culture medium containing 0.5% of polypeptone for 8 hours at 30°C, was added to each prepared culture medium, and culture was conducted for 42 hours at 30°C. After the culture, the cells were recovered by centrifuging, rinsed with methanol and lyophilized. The dried cells, after weighing, were agitated with chloroform for 48 hours at 50°C to extract a polymer. The chloroform extract was filtered, then

- 137 -

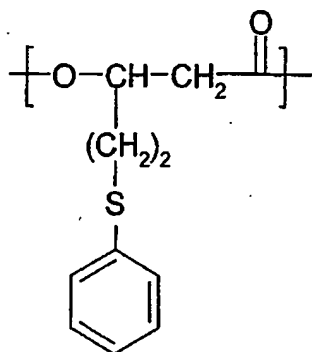
concentrated in an evaporator, and a solid precipitate formed with cold methanol was collected and dried under a reduced pressure to obtain a desired polymer. According to a weighing of the
5 obtained polymer, 890 mg (dry weight) of PHA were obtained in the present example.

An average molecular weight of the obtained PHA was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H,
10 solvent: chloroform, converted to polystyrene). As a result there were obtained a number-averaged molecular weight $M_n = 84000$ and a weight-averaged molecular weight $M_w = 169000$.

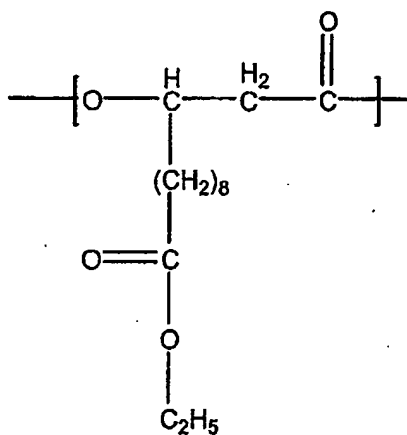
For specifying the structure of the obtained
15 PHA, a NMR analysis was conducted under same conditions as in Example 1.

As a result, there was confirmed a polyhydroxy alkanooate copolymer including, as monomer units, 3-hydroxy-5-(phenylsulfanyl)valeric acid represented by
20 the following chemical formula (58) by 80 mol%, three units of 3-hydroxy-11-ethoxycarbonylundecanoic acid represented by the following chemical formula (69), 3-hydroxy-9-ethoxycarbonylnonanoic acid represented by a chemical formula (70), and 3-hydroxy-7-
25 ethoxycarbonylheptanoic acid represented by a chemical formula (71) collectively by 14 mol%, and others (linear 3-hydroxyalkanoic acid of 4 to 12

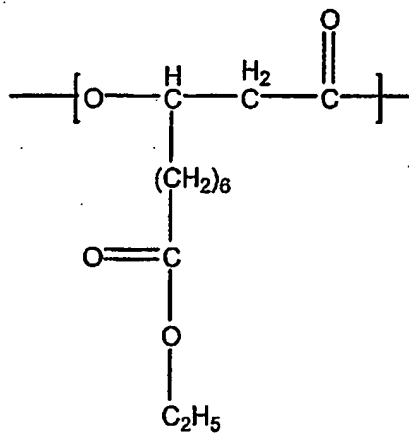
carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or 12 carbon atoms) by 6 mol%.



(58)

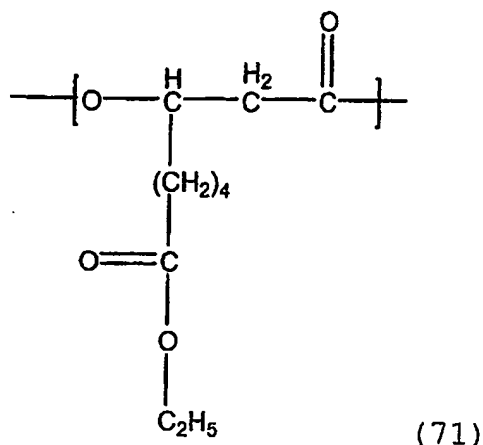


(69)



(70)

- 139 -



[Example 28]

There were prepared two 2000-ml shake flasks, and, in each, 0.5 wt.% of polypeptone (supplied by Wako Pure Chemicals Co.), 4 mmol/L of 5-benzoylvaleric acid, and 1 mmol/L of dodecanedioic acid monoethyl ester were dissolved in 1000 ml of an aforementioned M9 culture medium, which was placed in a 2000 ml shake flask, then sterilized in an autoclave and cooled to the room temperature. Then 5 ml of a culture liquid of *Pseudomonas cichorii* YN2 strain, shake cultured in advance in an M9 culture medium containing 0.5% of polypeptone for 8 hours at 30°C, was added to each prepared culture medium, and culture was conducted for 41 hours at 30°C. After the culture, the cells were recovered by centrifuging, rinsed with methanol and lyophilized. The dried cells, after weighing, were agitated with chloroform for 48 hours at 50°C to extract a polymer. The chloroform extract was filtered, then concentrated in

- 140 -

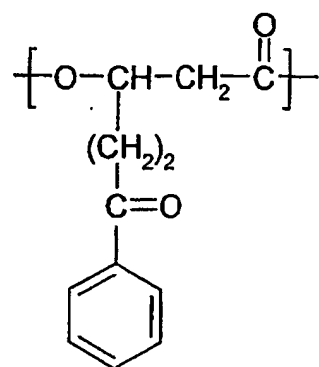
an evaporator, and a solid precipitate formed with cold methanol was collected and dried under a reduced pressure to obtain a desired polymer. According to a weighing of the obtained polymer, 450 mg (dry weight) of PHA were obtained in the present example.

An average molecular weight of the obtained PHA was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a result there were obtained a number-averaged molecular weight $M_n = 156000$ and a weight-averaged molecular weight $M_w = 325000$.

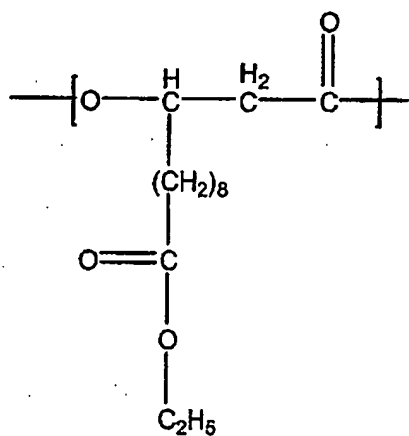
For specifying the structure of the obtained PHA, a NMR analysis was conducted under same conditions as in Example 1.

As a result, there was confirmed a polyhydroxy alkanooate copolymer including, as monomer units, 3-hydroxy-5-benzoylvaleric acid represented by the following chemical formula (62) by 69 mol%, three units of 3-hydroxy-11-ethoxycarbonylundecanoic acid represented by the following chemical formula (69), 3-hydroxy-9-ethoxycarbonylnonanoic acid represented by a chemical formula (70), and 3-hydroxy-7-ethoxycarbonylheptanoic acid represented by a chemical formula (71) collectively by 18 mol%, and others (linear 3-hydroxyalkanoic acid of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or

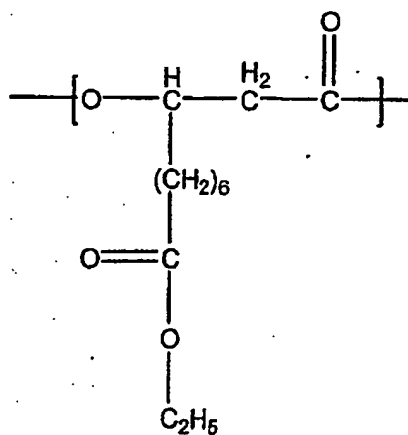
12 carbon atoms) by 13 mol%.



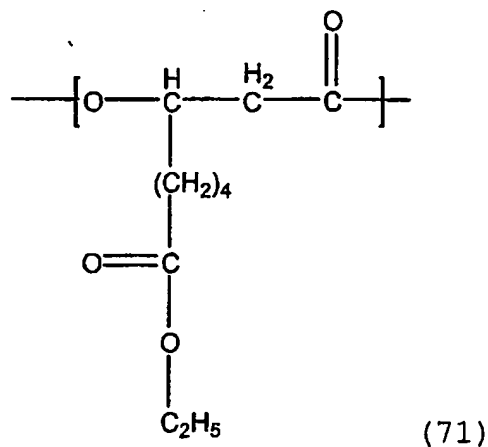
(62)



(69)



(70)



[Example 29]

There were prepared two 2000-ml shake flasks, and, in each, 0.5 wt.% of polypeptone (supplied by Wako Pure Chemicals Co.), 4 mmol/L of 5-(4-cyanophenoxy)valeric acid, and 1 mmol/L of sebacic acid monomethyl ester were dissolved in 1000 ml of an aforementioned M9 culture medium, which was placed in a 2000 ml shake flask, then sterilized in an autoclave and cooled to the room temperature. Then 5 ml of a culture liquid of *Pseudomonas cichorii* YN2 strain, shake cultured in advance in an M9 culture medium containing 0.5% of polypeptone for 8 hours at 30°C, was added to each prepared culture medium, and culture was conducted for 41 hours at 30°C. After the culture, the cells were recovered by centrifuging, rinsed with methanol and lyophilized. The dried cells, after weighing, were agitated with chloroform for 48 hours at 50°C to extract a polymer. The chloroform extract was filtered, then concentrated in

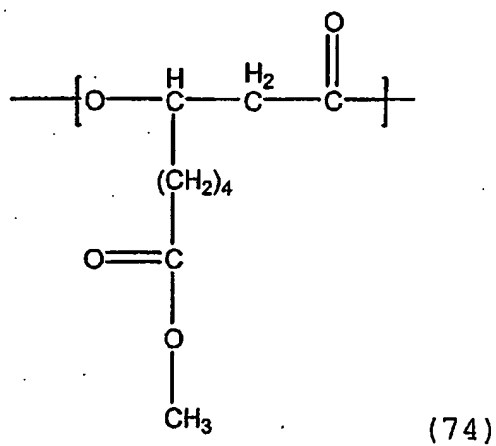
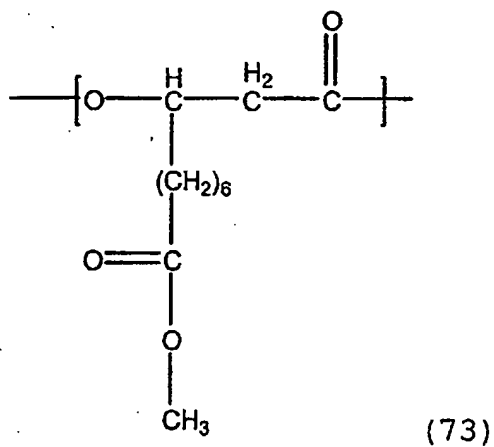
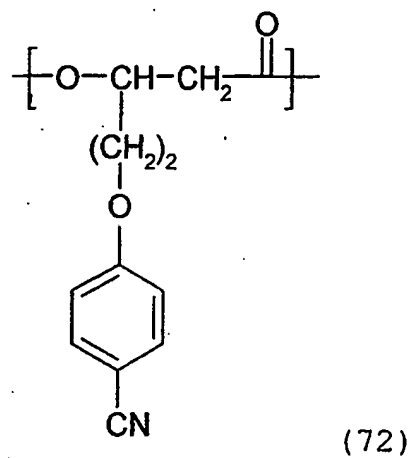
- 143 -

an evaporator, and a solid precipitate formed with cold methanol was collected and dried under a reduced pressure to obtain a desired polymer. According to a weighing of the obtained polymer, 450 mg (dry weight) of PHA were obtained in the present example.

An average molecular weight of the obtained PHA was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a result there were obtained a number-averaged molecular weight $M_n = 68000$ and a weight-averaged molecular weight $M_w = 129000$.

For specifying the structure of the obtained PHA, a NMR analysis was conducted under same conditions as in Example 1.

As a result, there was confirmed a polyhydroxy alkanooate copolymer including, as monomer units, 3-hydroxy-5-(4-cyanophenoxy)valeric acid represented by the following chemical formula (72) by 34 mol%, two units of 3-hydroxy-9-methoxycarbonylnonanoic acid represented by the following chemical formula (73) and 3-hydroxy-7-methoxycarbonylheptanoic acid represented by a chemical formula (74) collectively by 16 mol%, and others (linear 3-hydroxyalkanoic acid of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or 12 carbon atoms) by 50 mol%.



[Example 30]

5 There were prepared two 2000-ml shake flasks,
and, in each, 0.1 wt.% of n-nonanoic acid (supplied

- 145 -

by Kishida Kagaku), 4 mmol/L of 5-(4-nitrophenyl)valeric acid, and 1 mmol/L of sebacic acid monomethyl ester were dissolved in 1000 ml of an aforementioned M9 culture medium, which was placed in a 2000 ml shake flask, then sterilized in an autoclave and cooled to the room temperature. Then 5 ml of a culture liquid of *Pseudomonas cichorii* YN2 strain, shake cultured in advance in an M9 culture medium containing 0.5% of polypeptone for 8 hours at 30°C, was added to each prepared culture medium, and culture was conducted for 72 hours at 30°C. After the culture, the cells were recovered by centrifuging, rinsed with methanol and lyophilized. The dried cells, after weighing, were agitated with chloroform for 48 hours at 50°C to extract a polymer. The chloroform extract was filtered, then concentrated in an evaporator, and a solid precipitate formed with cold methanol was collected and dried under a reduced pressure to obtain a desired polymer. According to a weighing of the obtained polymer, 170 mg (dry weight) of PHA were obtained in the present example.

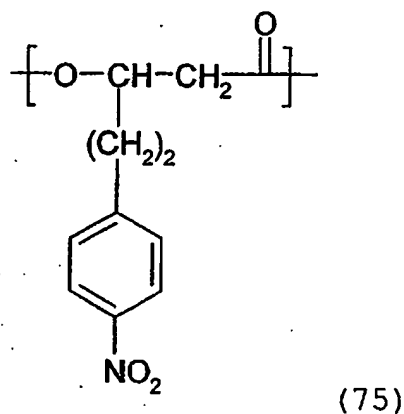
An average molecular weight of the obtained PHA was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a result there were obtained a number-averaged molecular weight $M_n = 59000$ and a weight-averaged

- 146 -

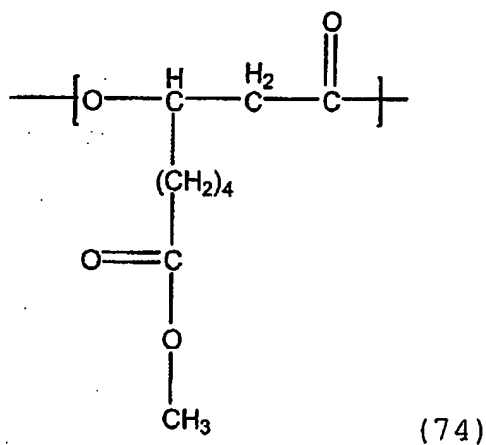
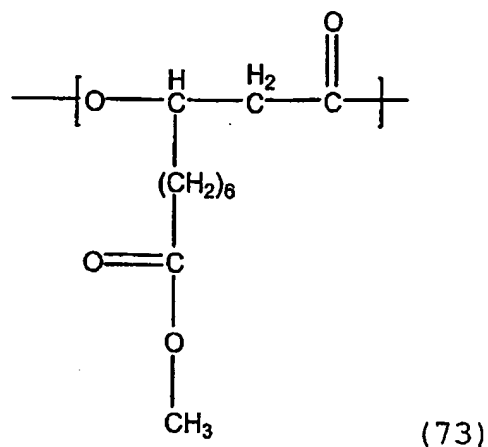
molecular weight Mw = 125000.

For specifying the structure of the obtained PHA, a NMR analysis was conducted under same conditions as in Example 1.

5 As a result, there was confirmed a polyhydroxy alkanate copolymer including, as monomer units, 3-hydroxy-5-(4-nitrophenyl)valeric acid represented by the following chemical formula (75) by 8 mol%, two units of 3-hydroxy-9-methoxycarbonylnonanoic acid
10 represented by the following chemical formula (73) and 3-hydroxy-7-methoxycarbonylheptanoic acid represented by a chemical formula (74) collectively by 18 mol%, and others (linear 3-hydroxyalkanoic acid of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid
15 with 10 or 12 carbon atoms) by 74 mol%.



- 147 -



[Example 31]

There were prepared two 2000-ml shake flasks,
 5 and, in each, 0.1 wt.% of n-nonanoic acid (supplied
 by Kishida Kagaku), 4 mmol/L of 5-
 [(phenylmethyl)oxy]valeric acid, and 1 mmol/L of
 sebacic acid monomethyl ester were dissolved in 1000
 ml of an aforementioned M9 culture medium, which was
 10 placed in a 2000 ml shake flask, then sterilized in
 an autoclave and cooled to the room temperature.
 Then 5 ml of a culture liquid of *Pseudomonas cichorii*
 YN2 strain, shake cultured in advance in an M9

- 148 -

culture medium containing 0.5% of polypeptone for 8 hours at 30°C, was added to each prepared culture medium, and culture was conducted for 40 hours at 30°C. After the culture, the cells were recovered by centrifuging, rinsed with methanol and lyophilized. The dried cells, after weighing, were agitated with chloroform for 48 hours at 50°C to extract a polymer. The chloroform extract was filtered, then concentrated in an evaporator, and a solid precipitate formed with cold methanol was collected and dried under a reduced pressure to obtain a desired polymer. According to a weighing of the obtained polymer, 330 mg (dry weight) of PHA were obtained in the present example.

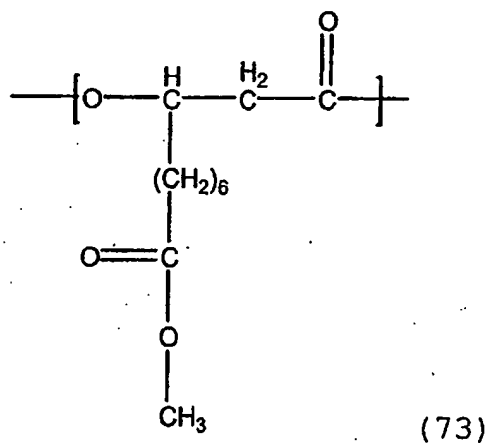
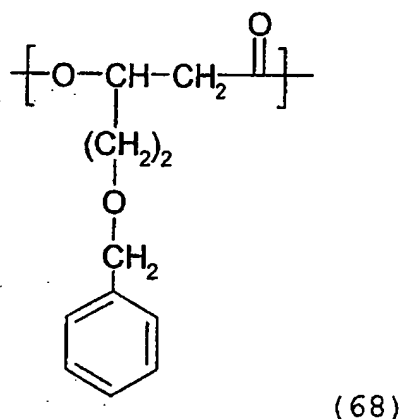
An average molecular weight of the obtained PHA was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a result there were obtained a number-averaged molecular weight $M_n = 79000$ and a weight-averaged molecular weight $M_w = 152000$.

For specifying the structure of the obtained PHA, a NMR analysis was conducted under same conditions as in Example 1.

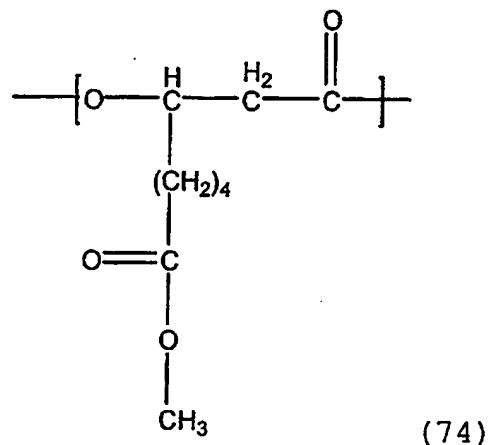
As a result, there was confirmed a polyhydroxy alkanoate copolymer including, as monomer units, 3-hydroxy-5-[(phenylmethyl)oxy]valeric acid represented

- 149 -

- by the following chemical formula (68) by 81 mol%,
 two units of 3-hydroxy-9-methoxycarbonylnonanoic acid
 represented by the following chemical formula (73)
 and 3-hydroxy-7-methoxycarbonylheptanoic acid
 5 represented by a chemical formula (74) collectively
 by 13 mol%, and others (linear 3-hydroxyalkanoic acid
 of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid
 with 10 or 12 carbon atoms) by 6 mol%.



- 150 -



[Example 32]

There were prepared two 2000-ml shake flasks, and, in each, 0.5 wt.% of polypeptone (supplied by

5 Wako Pure Chemical Co.), 4 mmol/L of 5-5-phenylvaleric acid, and 1 mmol/L of sebacic acid monomethyl ester were dissolved in 1000 ml of an aforementioned M9 culture medium, which was placed in a 2000 ml shake flask, then sterilized in an

10 autoclave and cooled to the room temperature. Then 5 ml of a culture liquid of *Pseudomonas cichorii* YN2 strain, shake cultured in advance in an M9 culture medium containing 0.5% of polypeptone for 8 hours at 30°C, was added to each prepared culture medium, and

15 culture was conducted for 40 hours at 30°C. After the culture, the cells were recovered by centrifuging, rinsed with methanol and lyophilized. The dried cells, after weighing, were agitated with chloroform for 48 hours at 50°C to extract a polymer. The

20 chloroform extract was filtered, then concentrated in

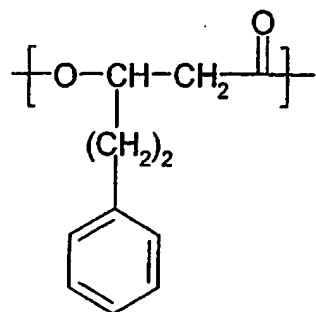
- 151 -

an evaporator, and a solid precipitate formed with cold methanol was collected and dried under a reduced pressure to obtain a desired polymer. According to a weighing of the obtained polymer, 1340 mg (dry weight) of PHA were obtained in the present example.

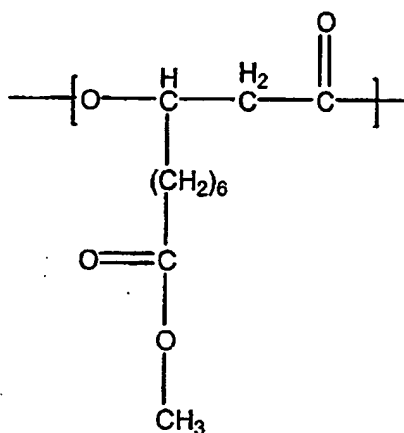
An average molecular weight of the obtained PHA was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a result there were obtained a number-averaged molecular weight $M_n = 81000$ and a weight-averaged molecular weight $M_w = 159000$.

For specifying the structure of the obtained PHA, a NMR analysis was conducted under same conditions as in Example 1.

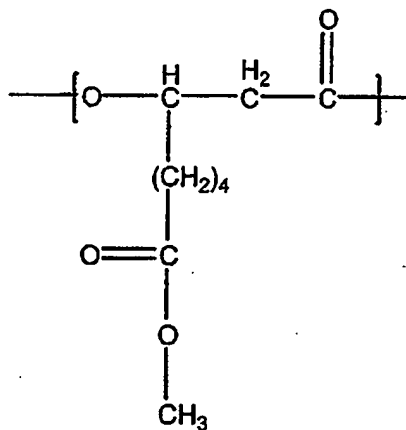
As a result, there was confirmed a polyhydroxy alkanooate copolymer including, as monomer units, 3-hydroxy-5-phenylvaleric acid represented by the following chemical formula (60) by 77 mol%, two units of 3-hydroxy-9-methoxycarbonylnonanoic acid represented by the following chemical formula (73) and 3-hydroxy-7-methoxycarbonylheptanoic acid represented by a chemical formula (74) collectively by 19 mol%, and others (linear 3-hydroxyalkanoic acid of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or 12 carbon atoms) by 4 mol%.



(60)



(73)



(74)

The polyhydroxy alkanoate thus obtained was
5 utilized in the following reaction.

The synthesized polyhydroxy alkanoate was
formed into a film, and 500 mg of such film was
placed on a Petri dish and was let to stand for 5

- 153 -

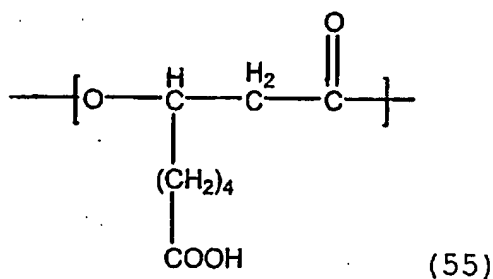
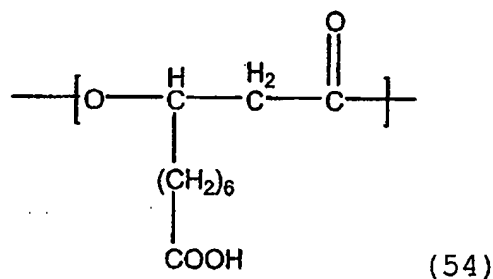
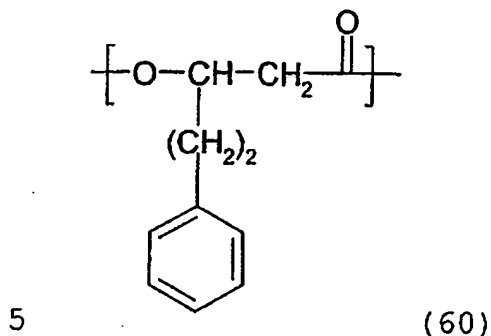
hours in 100 ml of a 0.1N aqueous solution of sodium hydroxide. After the reaction, the sodium hydroxide solution was removed, and the polymer was washed three times with 100 ml of distilled water. Then the
5 polymer was dissolved in 200 ml of ethyl acetate, and, after an addition of 100 ml of a 1.0N aqueous solution of hydrochloric acid, the solution was agitated for 1 hour at the room temperature. Then the polymer was extracted, washed with distilled
10 water and the solvent was distilled off to recover the polymer. Thereafter, a drying under a reduced pressure was carried out to obtain 350 mg of a desired polymer.

An average molecular weight of the obtained PHA
15 was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a result there were obtained a number-averaged molecular weight $M_n = 9500$ and a weight-averaged
20 molecular weight $M_w = 32000$.

For specifying the structure of the obtained PHA, a NMR analysis was conducted under same conditions as in Example 1.

As a result, there was confirmed a polyhydroxy
25 alkanoate copolymer including, as monomer units, 3-hydroxy-5-phenylvaleric acid represented by the following chemical formula (60), 3-hydroxy-9-

carboxynonanoic acid represented by the following chemical formula (54) and 3-hydroxy-7-carboxyheptanoic acid represented by a chemical formula (55).



Also a proportion of the units in the obtained polymer was calculated from a decrease in ester groups, and confirmed as 3-hydroxy-5-phenylvaleric acid by 78 mol%, 3-hydroxy-9-carboxynonanoic acid and 3-hydroxy-7-carboxylheptanoic acid collectively by 12 mol%, 3-hydroxy-9-methoxycarbonylnonanoic acid and 3-hydroxy-7-methoxycarbonylheptanoic acid

- 155 -

collectively by 6 mol%, and others (linear 3-hydroxyalkanoic acid of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or 12 carbon atoms) by 4 mol%.

5 [Example 33]

There were prepared two 2000-ml shake flasks, and, in each, 0.5 wt.% of polypeptone (supplied by Wako Pure Chemical Co.), 4 mmol/L of 5-phenoxyvaleric acid, and 1 mmol/L of sebacic acid monomethyl ester
10 were dissolved in 1000 ml of an aforementioned M9 culture medium, which was placed in a 2000 ml shake flask, then sterilized in an autoclave and cooled to the room temperature. Then 5 ml of a culture liquid of *Pseudomonas cichorii* YN2 strain, shake cultured in
15 advance in an M9 culture medium containing 0.5% of polypeptone for 8 hours at 30°C, was added to each prepared culture medium, and culture was conducted for 40 hours at 30°C. After the culture, the cells were recovered by centrifuging, rinsed with methanol
20 and lyophilized. The dried cells, after weighing, were agitated with chloroform for 48 hours at 50°C to extract a polymer. The chloroform extract was filtered, then concentrated in an evaporator, and a solid precipitate formed with cold methanol was
25 collected and dried under a reduced pressure to obtain a desired polymer. According to a weighing of the obtained polymer, 710 mg (dry weight) of PHA were

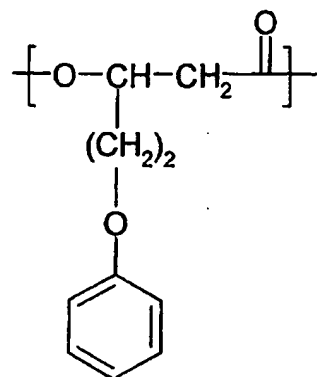
obtained in the present example.

An average molecular weight of the obtained PHA was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a result there were obtained a number-averaged molecular weight $M_n = 71000$ and a weight-averaged molecular weight $M_w = 148000$.

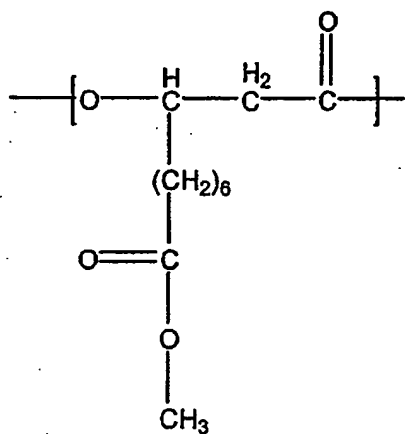
For specifying the structure of the obtained PHA, a NMR analysis was conducted under same conditions as in Example 1.

As a result, there was confirmed a polyhydroxy alkanoate copolymer including, as monomer units, 3-hydroxy-5-phenoxyvaleric acid represented by the following chemical formula (53) by 74 mol%, two units of 3-hydroxy-9-methoxycarbonylnonanoic acid represented by the following chemical formula (73) and 3-hydroxy-7-methoxycarbonylheptanoic acid represented by a chemical formula (74) collectively by 18 mol%, and others (linear 3-hydroxyalkanoic acid of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or 12 carbon atoms) by 8 mol%.

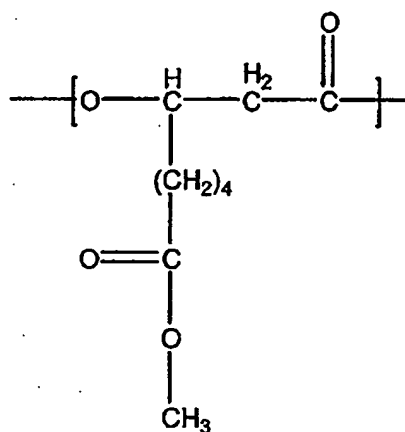
- 157 -



(53)



(73)



(74)

The polyhydroxy alkanoate thus obtained was
5 utilized in the following reaction.

The synthesized polyhydroxy alkanoate was
formed into a film, and 500 mg of such film was

- 158 -

placed on a Petri dish and was let to stand for 5 hours in 100 ml of a 0.1N aqueous solution of sodium hydroxide. After the reaction, the sodium hydroxide solution was removed, and the polymer was washed
5 three times with 100 ml of distilled water. Then the polymer was dissolved in 200 ml of ethyl acetate, and, after an addition of 100 ml of a 1.0N aqueous solution of hydrochloric acid, the solution was agitated for 1 hour at the room temperature. Then
10 the polymer was extracted, washed with distilled water and the solvent was distilled off to recover the polymer. Thereafter, a drying under a reduced pressure was carried out to obtain 370 mg of a desired polymer.

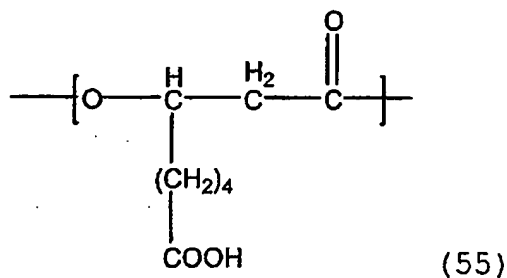
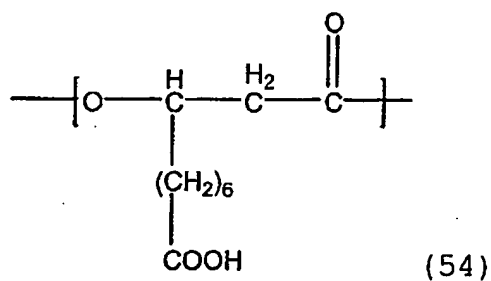
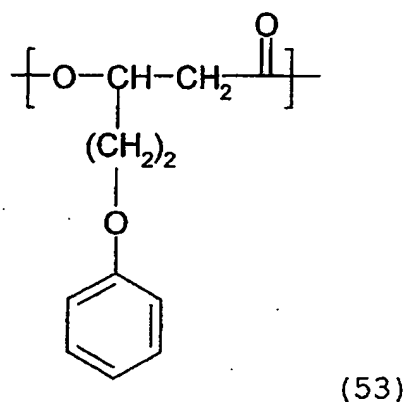
15 An average molecular weight of the obtained PHA was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). As a result there were obtained a number-averaged
20 molecular weight $M_n = 8700$ and a weight-averaged molecular weight $M_w = 30900$.

For specifying the structure of the obtained PHA, a NMR analysis was conducted under same conditions as in Example 1.

25 As a result, there was confirmed a polyhydroxy alkanoate copolymer including, as monomer units, 3-hydroxy-5-phenoxyvaleric acid represented by the

- 159 -

following chemical formula (53), 3-hydroxy-9-carboxynonanoic acid represented by the following chemical formula (54) and 3-hydroxy-7-carboxyheptanoic acid represented by a chemical
 5 formula (55).



Also a proportion of the units in the obtained
 10 polymer was calculated from a decrease in ester groups, and confirmed as 3-hydroxy-5-phenoxyvaleric acid by 73 mol%, 3-hydroxy-9-carboxynonanoic acid and 3-hydroxy-7-carboxyheptanoic acid collectively by 10

- 160 -

mol%, 3-hydroxy-9-methoxycarbonylnonanoic acid and 3-hydroxy-7-methoxycarbonylheptanoic acid collectively by 8 mol%,
and others (linear 3-hydroxyalkanoic acid of 4 to 12 carbon atoms and 3-hydroxyalk-5-enoic acid with 10 or 12 carbon atoms) by 9 mol%.

[Example 34]

In 1000 mL of an aforementioned M9 culture medium, there were added 0.5 wt.% of polypeptone
10 (supplied by Wako Pure Chemical Co.), and 5-phenylvaleric acid and sebacic acid monomethyl ester so as to obtain final concentrations of 4 and 1 mmol/L respectively, and the solution was placed in a 2000 ml shake flask, then sterilized in an autoclave
15 and cooled to the room temperature. Then 5 ml of a culture liquid of *Pseudomonas cichorii* YN2 strain, shake cultured in advance in an M9 culture medium containing 0.5% of polypeptone for 8 hours at 30°C, was added to each prepared culture medium, and
20 culture was conducted for 40 hours at 30°C. After the culture, the cells were recovered by centrifuging, rinsed with methanol and lyophilized. The dried cells, after weighing, were agitated with chloroform for 48 hours at 50°C to extract a polymer. The
25 chloroform extract was filtered, then concentrated in an evaporator, and a solid precipitate formed with cold methanol was collected and dried under a reduced

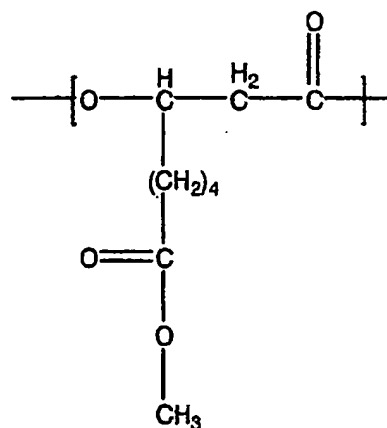
- 161 -

pressure to obtain a desired polymer.

A structure determination of the obtained polymer was carried out by a methynolysis-GC/MS method to be explained in the following. 5 mg of the polymer were dissolved in 2 mL of chloroform, then added with 2 mL of a 3% methanol solution of sulfuric acid and refluxed for 3.5 hours at 100°C. After the reaction, the reaction mixture was cooled to the room temperature and separated by adding 10 mL of deionized water under agitation. Then an organic layer was dehydrated with magnesium sulfate (anhydrous) and the reaction liquid was subjected to a measurement by a gas chromatography-mass spectrometer (GC/MS: Shimadzu QP-5050A, column: DB-WAXETR 0.32 mm x 30 m). An obtained total ion chromatogram (TIC) is shown in Fig. 5. There were observed three main peaks at 35.6, 38.0 and 45.8 minutes. A mass spectrum (MS) of the peak at 35.6 minutes is shown in Fig. 6; a MS of the peak at 38.0 minutes is shown in Fig. 7; and a MS of the peak at 45.8 minutes is shown in Fig. 8.

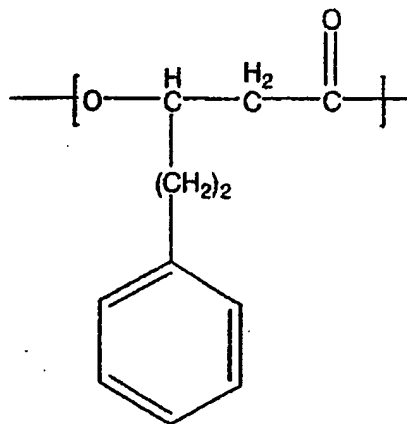
As a result, the peak at 35.6 minutes was derived from a unit shown in a chemical formula (80):

- 162 -



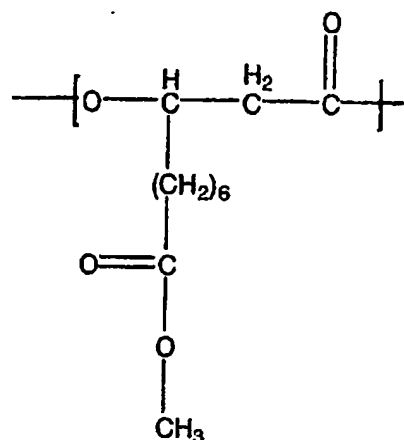
(80)

the peak at 38.0 minutes was derived from a unit shown in a chemical formula (81):



(81)

5 and the peak at 35.6 minutes was derived from a unit shown in a chemical formula (82):



(82)

Also proportions of the units, calculated from ratios of the peak areas of TIC, were 12.0%, 77.7% and 6.7% respectively.

- 5 The molecular weight of the polymer was measured by gel permeation chromatography (GPC: Toso HLC-8220 GPC, column: Toso TSK-GEL Super HM-H, solvent: chloroform, converted to polystyrene). Table 1 shows weights of the obtained cells and the
- 10 obtained polymer, a polymer weight ratio per cell, The molecular weight and The molecular weight distribution of the obtained polymer.

Table 1

CDW (mg/L)	PDW (mg/L)	P/C %	Mn ($\times 10^4$)	Mw ($\times 10^4$)	Mw/Mn
1358	671	49.4	8.1	15.9	2.0

- CDW: cell dry weight, PDW: polymer dry weight,
- 15 P/C: cell dry weight/polymer dry weight, Mn: number-averaged molecular weight,
- Mw: weight-averaged molecular weight,

- 164 -

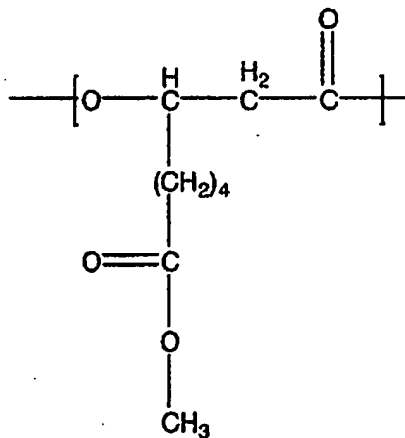
Mw/Mn: molecular weight distribution.

[Example 35]

A desired polymer was obtained in the same manner as in Example 34, except that the YN2 strain employed in Example 34 was replaced by *Pseudomonas jessenii* P161 strain.

A structure determination of the obtained polymer carried out by a methanolysis-GC/MS method as in Example 34 confirmed that the polymer was a polyhydroxy alkanoate copolymer constituted of units represented by chemical formulas (80), (81) and (82):

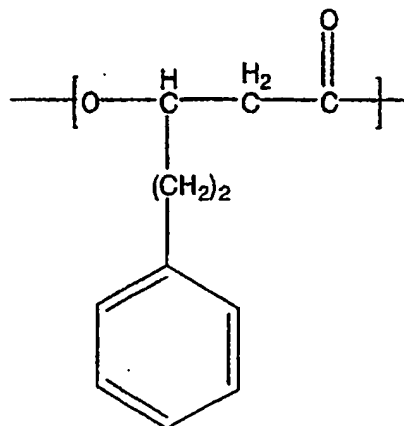
chemical formula (80):



(80)

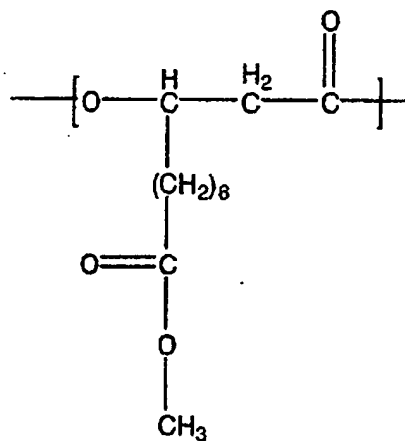
chemical formula (81):

- 165 -



(81)

chemical formula (82):



(82)

5 and the proportions of the units, calculated from the ratios of the peak areas of TIC, were 15.5%, 75.2% and 9.3% respectively.

The molecular weight of the polymer was measured by gel permeation chromatography as in
10 Example 34.

Table 2 shows weights of the obtained cells and

the obtained polymer, a polymer weight ratio per cell, the molecular weight and the molecular weight distribution of the obtained polymer.

Table 2

CDW (mg/L)	PDW (mg/L)	P/C %	Mn ($\times 10^4$)	Mw ($\times 10^4$)	Mw/Mn
821	271	33.0	6.6	13.9	2.1

- 5 CDW: cell dry weight, PDW: polymer dry weight,
P/C: cell dry weight/polymer dry weight, Mn: number-
averaged molecular weight,
Mw: weight-averaged molecular weight,
Mw/Mn: molecular weight distribution.

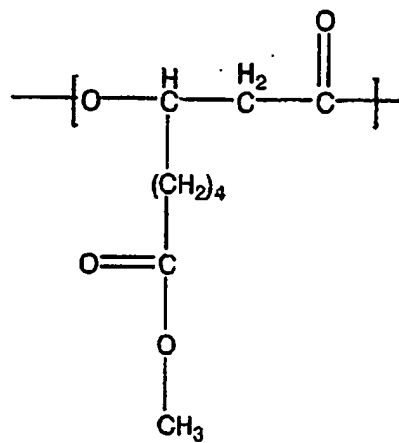
10 [Example 36]

A desired polymer was obtained in the same manner as in Example 34, except that the YN2 strain employed in Example 34 was replaced by *Pseudomonas cichorii* H45 strain and polypeptone was replaced by
15 yeast extract (DIFCO).

A structure determination of the obtained polymer carried out by a methanolysis-GC/MS method as in Example 34 confirmed that the polymer was a polyhydroxy alkanoate copolymer constituted of units
20 represented by chemical formulas (80), (81) and (82):

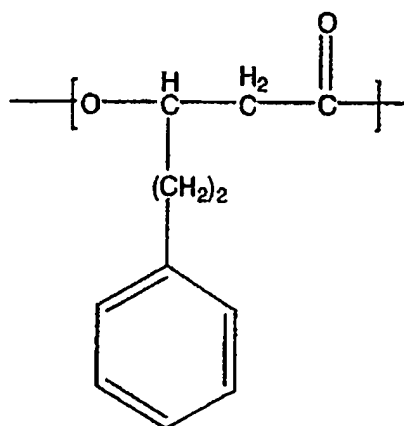
chemical formula (80):

- 167 -



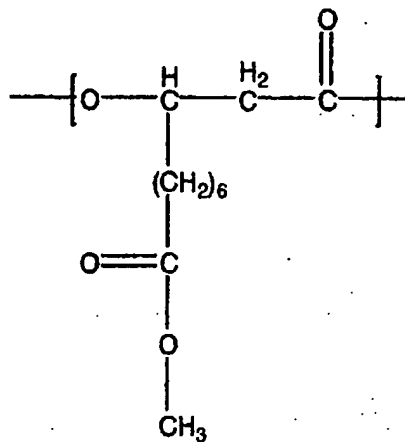
(80)

chemical formula (81):



(81)

chemical formula (82):



(82)

- 168 -

and the proportions of the units calculated from the ratios of the peak areas of TIC were 16.1%, 72.3% and 11.6% respectively.

The molecular weight of the polymer was measured by gel permeation chromatography as in Example 34.

Table 3 shows weights of the obtained cells and the obtained polymer, polymer weight ratio per cell, the molecular weight and the molecular weight distribution of the obtained polymer.

Table 3

CDW (mg/L)	PDW (mg/L)	P/C %	Mn ($\times 10^4$)	Mw ($\times 10^4$)	Mw/Mn
779	230	29.5	7.2	14.9	2.1

CDW: cell dry weight, PDW: polymer dry weight,

P/C: cell dry weight/polymer dry weight, Mn: number-averaged molecular weight,

Mw: weight-averaged molecular weight,

Mw/Mn: molecular weight distribution.

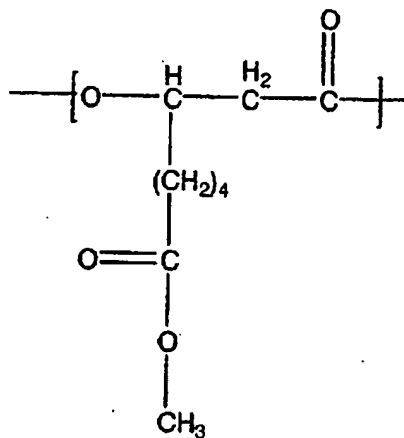
[Example 37]

A desired polymer was obtained in the same manner as in Example 34, except that the YN2 strain employed in Example 34 was replaced by *Pseudomonas putida* P91 strain and polypeptone was replaced by nonanoic acid (Kishida Kagaku, concentration: 0.1%).

A structure determination of the obtained polymer carried out by a methanolysis-GC/MS method as in Example 34 confirmed that the polymer was a

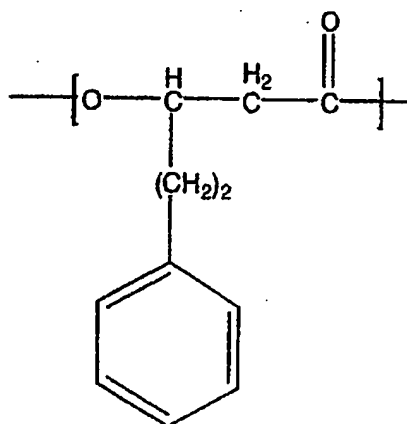
polyhydroxy alkanoate copolymer constituted of units represented by chemical formulas (80), (81), (82), (83), (84) and (85):

chemical formula (80):



(80)

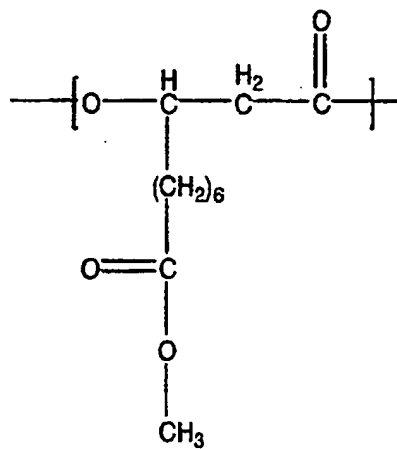
chemical formula (81):



(81)

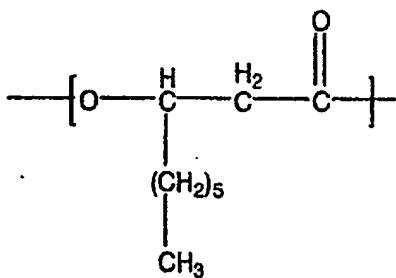
chemical formula (82)

- 170 -



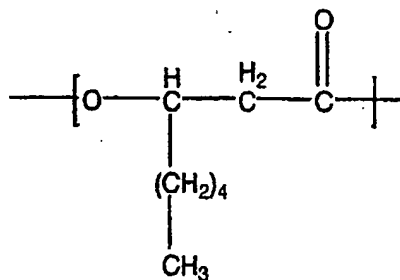
(82)

chemical formula (83):



(83)

chemical formula (84):

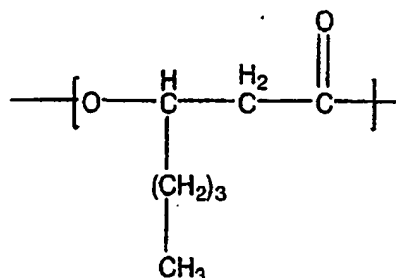


(84)

5

chemical formula (85):

- 171 -



(85)

and the proportions of the units calculated from the ratios of the peak areas of TIC were 5.1%, 52.1%, 6.6%, 11.3%, 4.9% and 20.0% respectively.

5 The molecular weight of the polymer was measured by gel permeation chromatography as in Example 34.

Table 4 shows weights of the obtained cells and the obtained polymer, a polymer weight ratio per cell,
10 a molecular weight and a molecular weight distribution of the obtained polymer.

Table 4

CDW (mg/L)	PDW (mg/L)	P/C %	Mn ($\times 10^4$)	Mw ($\times 10^4$)	Mw/Mn
528	110	20.8	8.2	16.9	2.1

CDW: cell dry weight, PDW: polymer dry weight,

P/C: cell dry weight/polymer dry weight, Mn: number-
15 averaged molecular weight,

Mw: weight-averaged molecular weight,

Mw/Mn: molecular weight distribution.

[Example 38]

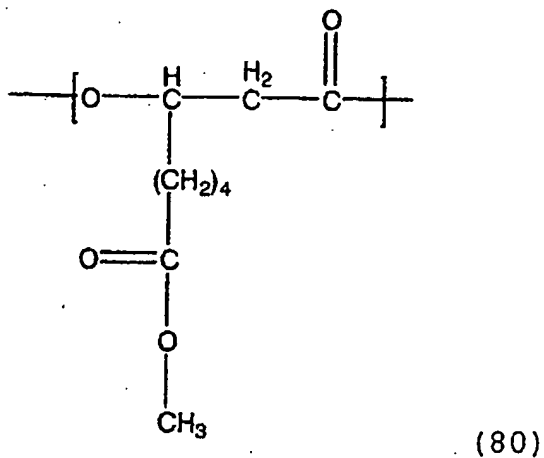
A desired polymer was obtained in the same

- 172 -

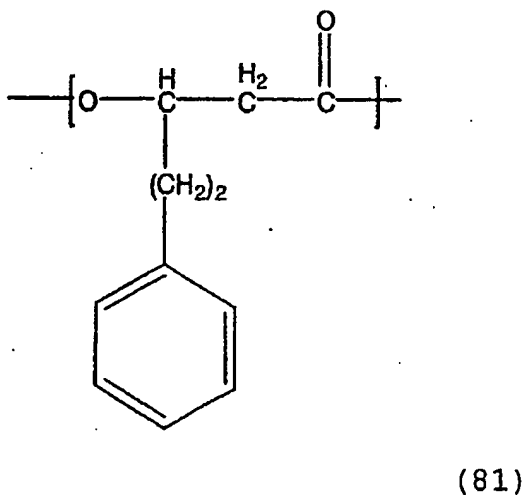
manner as in Example 34, except that polypeptone was replaced by D-glucose (Kishida Kagaku).

A structure determination of the obtained polymer carried out by a methanolysis-GC/MS method as in Example 34 confirmed that the polymer was a polyhydroxy alkanoate copolymer constituted of units represented by chemical formulas (80), (81) and (82):

chemical formula (80):

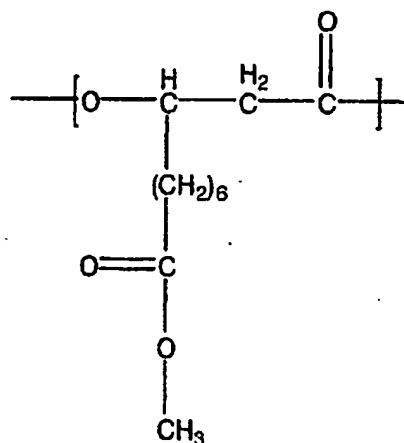


10 chemical formula (81):



chemical formula (82):

- 173 -



(82)

and the proportions of the units calculated from the ratios of the peak areas of TIC were 13.1%, 80.3% and 6.6% respectively.

5 The molecular weight of the polymer was measured by gel permeation chromatography as in Example 34.

Table 5 shows weights of the obtained cells and the obtained polymer, a polymer weight ratio per cell, a molecular weight and a molecular weight distribution of the obtained polymer.

Table 5

CDW (mg/L)	PDW (mg/L)	P/C %	Mn ($\times 10^4$)	Mw ($\times 10^4$)	Mw/Mn
910	425	46.7	7.9	15.4	1.9

CDW: cell dry weight, PDW: polymer dry weight,

P/C: cell dry weight/polymer dry weight, Mn: number-

15 averaged molecular weight,

Mw: weight-averaged molecular weight,

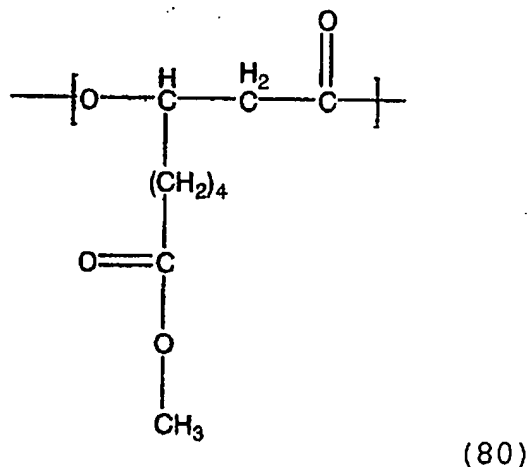
Mw/Mn: molecular weight distribution.

[Example 39]

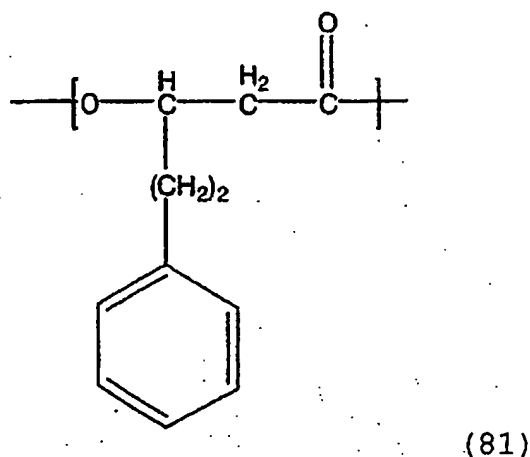
A desired polymer was obtained in the same manner as in Example 34, except that polypeptone was replaced by sodium piruvate (Kishida Kagaku).

5. A structure determination of the obtained polymer carried out by a methanolysis-GC/MS method as in Example 34 confirmed that the polymer was a polyhydroxy alkanoate copolymer constituted of units represented by chemical formulas (80), (81) and (82):

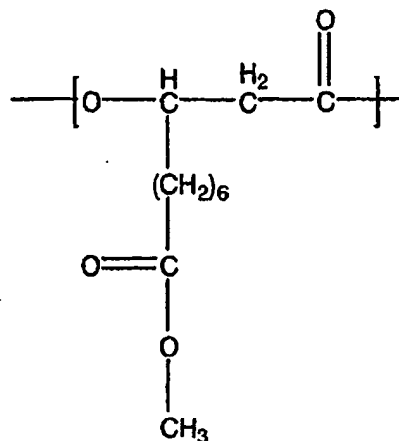
10 chemical formula (80):



chemical formula (81):



chemical formula (82):



(82)

and the proportions of the units calculated from the ratios of the peak areas of TIC were 11.9%, 82.2% and 5.9% respectively.

The molecular weight of the polymer was measured by gel permeation chromatography as in Example 34.

Table 6 shows weights of the obtained cells and the obtained polymer, a polymer weight ratio per cell, a molecular weight and a molecular weight distribution of the obtained polymer.

Table 6

CDW (mg/L)	PDW (mg/L)	P/C %	Mn ($\times 10^4$)	Mw ($\times 10^4$)	Mw/Mn
1120	585	52.2	8.0	15.9	2.0

CDW: cell dry weight, PDW: polymer dry weight,

P/C: cell dry weight/polymer dry weight, Mn: number-averaged molecular weight,

Mw: weight-averaged molecular weight,

- 176 -

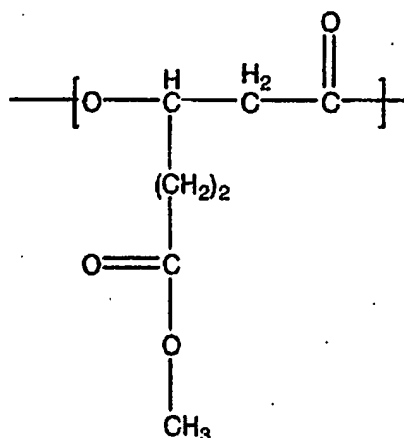
Mw/Mn: molecular weight distribution.

[Example 40]

A desired polymer was obtained in the same manner as in Example 34, except that polypeptone employed in Example 34 was replaced by sodium L-glutamate (Kishida Kagaku) and sebacic acid monomethyl ester, which is one of substrates for polymer synthesis was replaced by suberic acid monomethyl ester.

10 A structure determination of the obtained polymer carried out by a methanolysis-GC/MS method as in Example 34 confirmed that the polymer was a polyhydroxy alkanoate copolymer constituted of units represented by chemical formulas (86), (80) and (81):

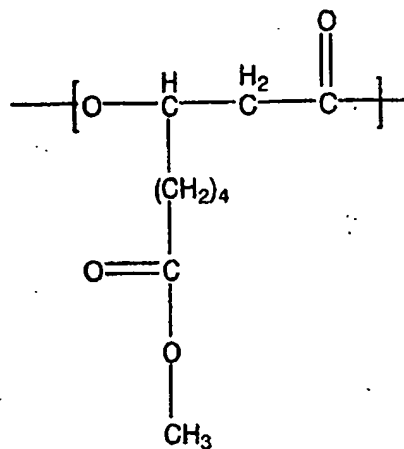
15 chemical formula (86):



(86)

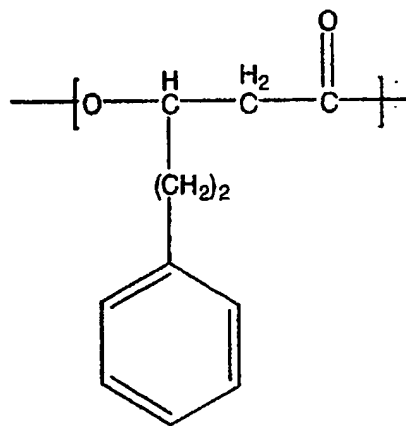
chemical formula (80):

- 177 -



(80)

chemical formula (81):



(81)

and the proportions of the units calculated from the ratios of the peak areas of TIC were 8.2%, 84.2% and 8.6% respectively.

The molecular weight of the polymer was measured by gel permeation chromatography as in Example 34.

Table 7 shows weights of the obtained cells and the obtained polymer, a polymer weight ratio per cell, a molecular weight and a molecular weight distribution of the obtained polymer.

5 Table 7

CDW (mg/L)	PDW (mg/L)	P/C %	Mn ($\times 10^4$)	Mw ($\times 10^4$)	Mw/Mn
985	440	44.7	7.8	14.6	1.9

CDW: cell dry weight, PDW: polymer dry weight,

P/C: cell dry weight/polymer dry weight, Mn: number-averaged molecular weight,

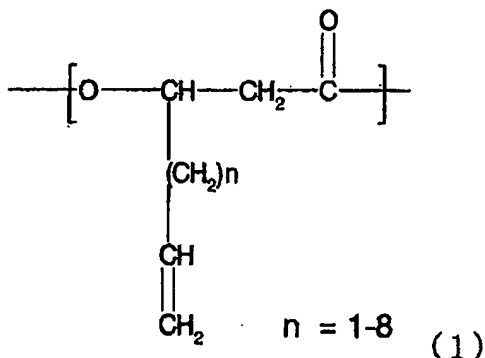
Mw: weight-averaged molecular weight,

10 Mw/Mn: molecular weight distribution.

- 179 -

CLAIMS

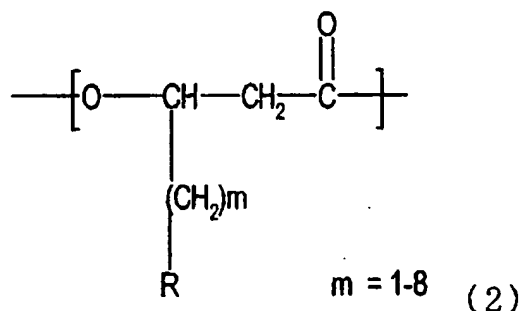
1. A polyhydroxy alkanooate copolymer characterized in including at least a 3-hydroxy- ω -alkenoic acid unit represented by a chemical formula (1) in a molecule, and simultaneously at least a 3-hydroxy- ω -alkanoic acid unit represented by a chemical formula (2) or a 3-hydroxy- ω -cyclohexylalkanoic acid unit represented by a chemical formula (3) in the molecule:
- 10 [Chemical Formula (1)]



- in which n represents an integer selected within a range indicated in the chemical formula; and in case plural units are present, n is the same or different for each unit;
- 15

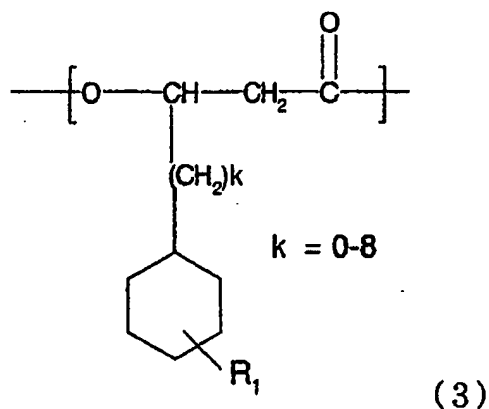
[Chemical Formula (2)]

- 180 -



in which m represents an integer selected within a range indicated in the chemical formula; R represents a residue having any of a phenyl structure or a thienyl structure; and in case plural units are present, m and R are the same or different for each unit;

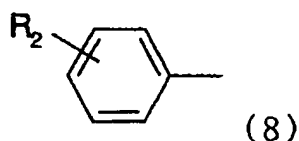
[Chemical Formula (3)]



in which R₁ being a substituent on a cyclohexyl group represents a hydrogen atom, a CN group, a NO₂ group, a halogen atom, a CH₃ group, a C₂H₅ group, a C₃H₇ group, a CF₃ group, a C₂F₅ group, or a C₃F₇ group; k represents an integer selected within a range indicated in the chemical formula; and in case plural units are present, R₁ and k may be the same or

different for each unit.

2. The polyhydroxy alkanoate copolymer according to claim 1, wherein R in the chemical
 5 formula (2) represents a residue having a phenyl structure or a thienyl structure selected from the group consisting of chemical formulas (8), (9), (10), (11), (12), (13), (14), (15), (16), (17) and (18):
 the chemical formula (8):

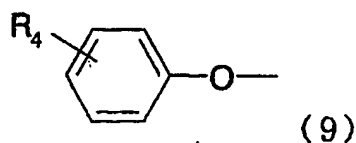


10

represents a group of non-substituted or substituted phenyl groups in which R_2 , a substituent on an aromatic ring and represents an H atom, represents a halogen atom, a CN group, a NO_2 group, a CH_3 group, a
 15 C_2H_5 group, a C_3H_7 group, a $\text{CH}=\text{CH}_2$ group, a COOR_3 group (R_3 represents an H atom, a Na atom or a K atom), a CF_3 group, a C_2F_5 group, or a C_3F_7 group; and in case plural units are present, R_2 is the same or different for each unit;

20

the chemical formula (9):

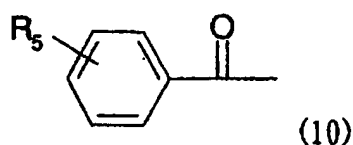


represents a group of non-substituted or substituted phenoxy groups in which R_4 represents a substituent on

- 182 -

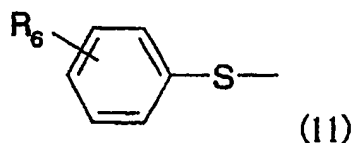
an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO₂ group, a CH₃ group, a C₂H₅ group, a C₃H₇ group, a SCH₃ group, a CF₃ group, a C₂F₅ group, or a C₃F₇ group; and in case plural units are
5 present, R₄ may be the same or different for each unit;

the chemical formula (10):



represents a group of non-substituted or substituted
10 benzoyl groups in which R₅ represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO₂ group, a CH₃ group, a C₂H₅ group, a C₃H₇ group, a CF₃ group, a C₂F₅ group, or a C₃F₇ group; and in case plural units are present, R₅
15 may be the same or different for each unit;

the chemical formula (11)

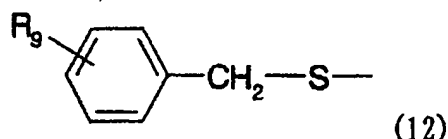


represents a group of substituted or non-substituted
phenylsulfanyl groups in which R₆ represents a
20 substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO₂ group, a COOR₇

- 183 -

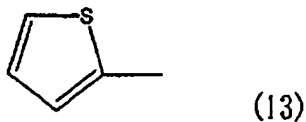
group, a SO_2R_8 group (R_7 represents either one of H, Na, K, CH_3 and C_2H_5 ; and R_8 represents either one of OH, ONa, OK, a halogen atom, OCH_3 and OC_2H_5), a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $(\text{CH}_3)_2\text{-CH}$ group or
 5 a $(\text{CH}_3)_3\text{-C}$ group; and in case plural units are present, R_6 may be the same or different for each unit;

the chemical formula (12):



represents a group of substituted or non-substituted
 10 (phenylmethyl)sulfanyl groups in which R_9 represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a COOR_{10} group, a SO_2R_{11} group (R_{10} represents either one of H, Na, K, CH_3 and C_2H_5 ; and R_{11} represents either
 15 one of OH, ONa, OK, a halogen atom, OCH_3 and OC_2H_5), a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $(\text{CH}_3)_2\text{-CH}$ group or a $(\text{CH}_3)_3\text{-C}$ group; and in case plural units are present, R_9 may be the same or different for each unit;

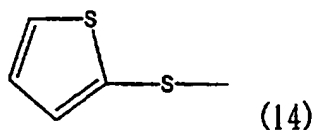
20 the chemical formula (13):



represents a 2-thienyl group;

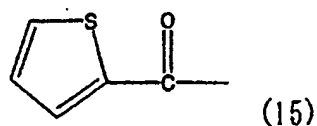
the chemical formula (14)

- 184 -



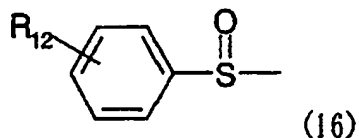
represents a 2-thienylsulfanyl group;

the chemical formula (15):



5 represents a 2-thienylcarbonyl group;

the chemical formula (16):



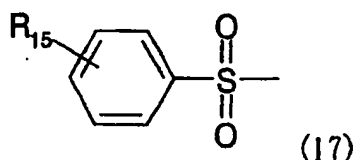
represents a group of substituted or non-substituted phenylsulfinyl groups in which R_{12} represents a

10 substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a COOR_{13} group, a SO_2R_{14} group (R_{13} represents either one of H, Na, K, CH_3 and C_2H_5 ; and R_{14} represents either one of OH, ONa, OK, a halogen atom, OCH_3 and OC_2H_5), a

15 CH_3 group, a C_2H_5 group, a C_3H_7 group, a $(\text{CH}_3)_2\text{-CH}$ group or a $(\text{CH}_3)_3\text{-C}$ group; and in case plural units are present, R_{12} may be the same or different for each unit;

the chemical formula (17):

- 185 -

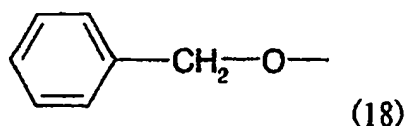


represents a group of substituted or non-substituted phenylsulfonyl groups in which R_{15} represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a

5 COOR₁₆ group, a SO₂R₁₇ group (R_{16} represents either one of H, Na, K, CH₃ and C₂H₅; and R_{17} represents either one of OH, ONa, OK, a halogen atom, OCH₃ and OC₂H₅), a CH₃ group, a C₂H₅ group, a C₃H₇ group, a (CH₃)₂-CH

10 group or a (CH₃)₃-C group; and in case plural units are present, R_{15} may be the same or different for each unit; and

the chemical formula (18):



15 represents a (phenylmethyl)oxy group.

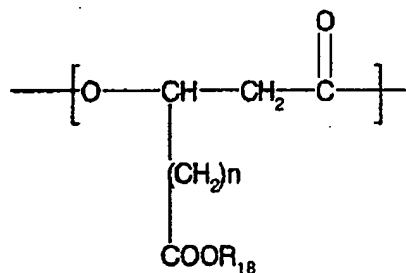
3. The polyhydroxy alkanoate copolymer according to claim 1, wherein a number-averaged molecular weight is within a range from 1000 to

20 1000000.

4. A polyhydroxy alkanoate copolymer characterized in including at least a 3-hydroxy- ω -

- 186 -

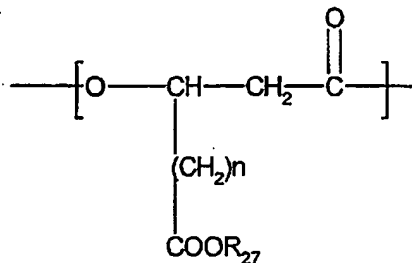
carboxyalkanoic acid unit represented by a chemical formula (19) or 3-hydroxy- ω -alkoxycarbonylalkanoic acid unit represented by a chemical formula (32) in a molecule, and simultaneously at least a 3-hydroxy- ω -alkanoic acid unit represented by a chemical formula (2) or a 3-hydroxy- ω -cyclohexylalkanoic acid unit represented by a chemical formula (3) in the molecule, [Chemical Formula (19)]



$$n = 1-8 \quad (19)$$

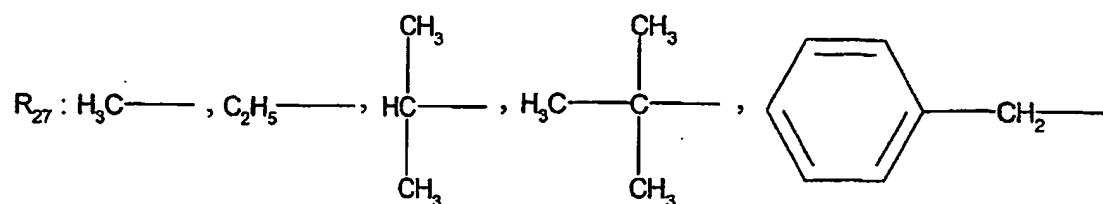
10 in which n represents an integer selected within a range indicated in the chemical formula; R_{18} represents an H atom, a Na atom or a K atom: and in case plural units are present, n and R_{18} may be the same or different for each unit; and

15 [Chemical Formula (32)]



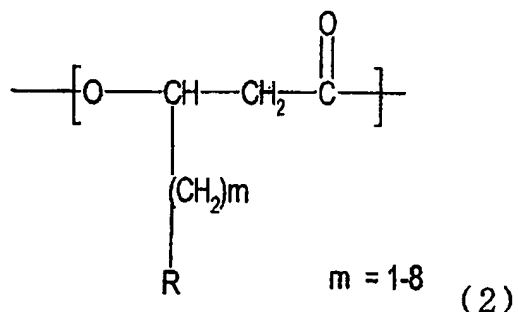
$$n = 1-8 \quad (32)$$

- 187 -



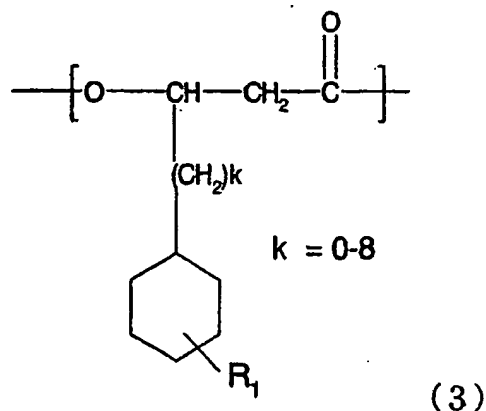
in which n represents an integer selected within a range indicated in the chemical formula; R_{27} represents any of residues indicated in the chemical formula; and in case plural units are present, n and R_{27} may be the same or different for each unit

[Chemical Formula (2)]



in which m represents an integer selected within a range indicated in the chemical formula; R includes a residue having any of a phenyl structure or a thienyl structure; and in case plural units are present, m and R may be the same or different for each unit; and [Chemical Formula (3)]

- 188 -

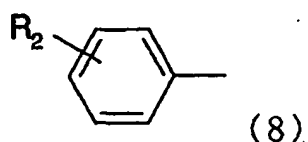


in which R_1 represents a substituent on a cyclohexyl group and represents an H atom, a CN group, a NO_2 group, a halogen atom, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a CF_3 group, a C_2F_5 group, or a C_3F_7 group; k represents an integer selected within a range indicated in the chemical formula; and in case plural units are present, R_1 and k are the same or different for each unit.

10

5. The polyhydroxy alkanoate copolymer according to claim 4, wherein R in the chemical formula (2), represents a residue having a phenyl structure or a thienyl structure selected from chemical formulas (8), (9), (10), (11), (12), (13), (14), (15), (16), (17), and (18):

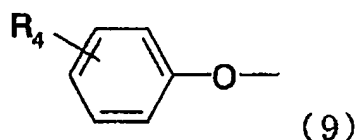
the chemical formula (8):



- 189 -

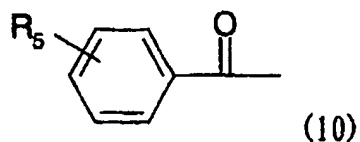
represents a group of non-substituted or substituted phenyl groups in which R_2 represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $\text{CH}=\text{CH}_2$ group, a COOR_3 group (R_3 representing an H atom, a Na atom or a K atom), a CF_3 group, a C_2F_5 group, or a C_3F_7 group; and in case plural units are present, R_2 is the same or different for each unit;

the chemical formula (9):



represents a group of non-substituted or substituted phenoxy groups in which R_4 represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a SCH_3 group, a CF_3 group, a C_2F_5 group, or a C_3F_7 group; and in case plural units are present, R_4 is the same or different for each unit;

the chemical formula (10):

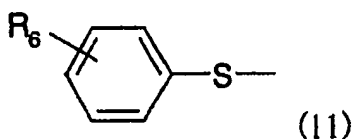


represents a group of non-substituted or substituted

- 190 -

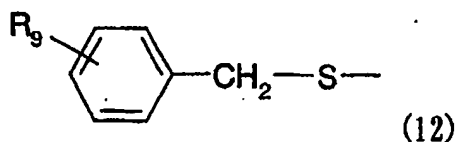
benzoyl groups in which R_5 represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a CF_3 group, a C_2F_5 group, or a
 5 C_3F_7 group; and in case plural units are present, R_5 is the same or different for each unit;

the chemical formula (11):



represents a group of substituted or non-substituted
 10 phenylsulfanyl groups in which R_6 represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a COOR_7 group, a SO_2R_8 group (R_7 represents either one of H, Na, K, CH_3 and C_2H_5 ; and R_8 represents either one of
 15 OH, ONa, OK, a halogen atom, OCH_3 and OC_2H_5), a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $(\text{CH}_3)_2\text{-CH}$ group or a $(\text{CH}_3)_3\text{-C}$ group; and in case plural units are present, R_6 is the same or different for each unit;

the chemical formula (12):



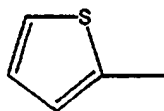
20

represents a group of substituted or non-substituted

- 191 -

(phenylmethyl)sulfanyl groups in which R_9 represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a COOR_{10} group, a SO_2R_{11} group (R_{10} represents either one
5 of H, Na, K, CH_3 and C_2H_5 ; and R_{11} represents either one of OH, ONa, OK, a halogen atom, OCH_3 and OC_2H_5), a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $(\text{CH}_3)_2\text{-CH}$ group or a $(\text{CH}_3)_3\text{-C}$ group; and in case plural units are present, R_9 is the same or different for each
10 unit;

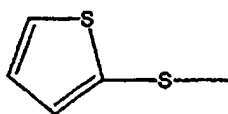
the chemical formula (13):



(13)

represents a 2-thienyl group;

the chemical formula (14):

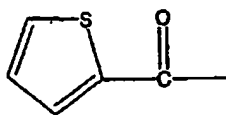


(14)

15

represents a 2-thienylsulfanyl group;

the chemical formula (15):



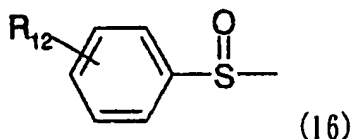
(15)

represents a 2-thienylcarbonyl group;

20

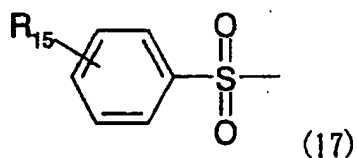
the chemical formula (16):

- 192 -



represents a group of substituted or non-substituted phenylsulfinyl groups in which R_{12} represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a COOR_{13} group, a SO_2R_{14} group (R_{13} represents either one of H, Na, K, CH_3 and C_2H_5 ; and R_{14} represents either one of OH, ONa, OK, a halogen atom, OCH_3 and OC_2H_5), a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $(\text{CH}_3)_2\text{-CH}$ group or a $(\text{CH}_3)_3\text{-C}$ group; and in case plural units are present, R_{12} is the same or different for each unit;

the chemical formula (17):

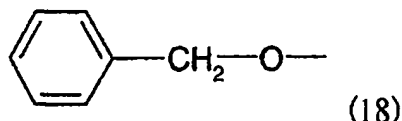


represents a group of substituted or non-substituted phenylsulfonyl groups in which R_{15} represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a COOR_{16} group, a SO_2R_{17} group (R_{16} represents either one of H, Na, K, CH_3 and C_2H_5 ; and R_{17} represents either one of OH, ONa, OK, a halogen atom, OCH_3 and OC_2H_5), a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $(\text{CH}_3)_2\text{-CH}$ group or a $(\text{CH}_3)_3\text{-C}$ group; and in case plural units

- 193 -

are present, R_{15} is the same or different for each unit; and

the chemical formula (18):



5 represents a (phenylmethyl)oxy group.

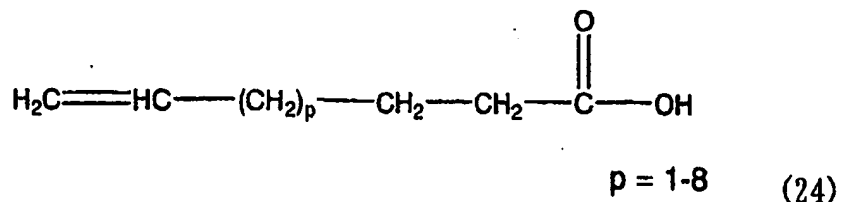
6. The polyhydroxy alkanoate copolymer according to claim 4, wherein a number-averaged molecular weight is within a range from 1000 to
10 1000000.

7. A method for producing a polyhydroxy alkanoate copolymer characterized in including a biosynthesis by a microorganism having an ability of
15 producing a polyhydroxy alkanoate copolymer including at least a 3-hydroxy- ω -alkenoic acid unit represented by a chemical formula (1) in a molecule, and simultaneously at least a 3-hydroxy- ω -alkanoic acid unit represented by a chemical formula (2) or a 3-
20 hydroxy- ω -cyclohexylalkanoic acid unit represented by a chemical formula (3) in the molecule, from at least an ω -alkenoic acid represented by a chemical formula (24) and at least a compound represented by a chemical formula (25) or at least an ω -
25 cyclohexylalkanoic acid represented by a chemical

- 194 -

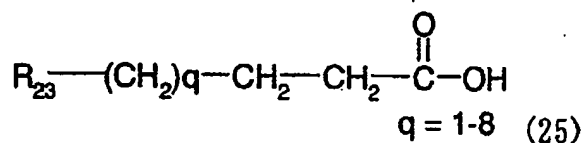
formula (26) as starting materials:

[Chemical Formula (24)]



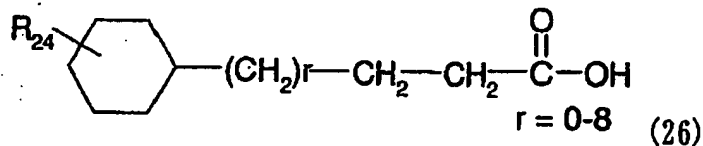
in which p represents an integer selected within a
5 range indicated in the chemical formula;

[Chemical Formula (25)]



in which q represents an integer selected within a
range indicated in the chemical formula; and R_{23}
10 includes a residue having a phenyl structure or a
thienyl structure;

[Chemical Formula (26)]

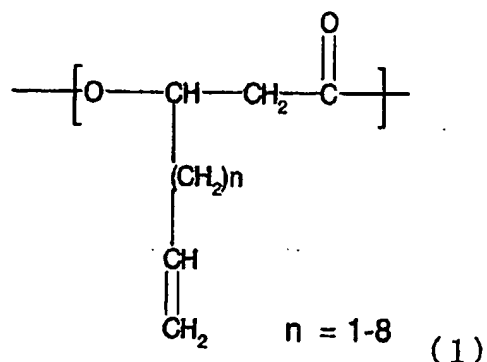


in which R_{24} represents a substituent on a cyclohexyl
15 group and represents an H atom, a CN group, a NO_2
group, a halogen atom, a CH_3 group, a C_2H_5 group, a
 C_3H_7 group, a CF_3 group, a C_2F_5 group, or a C_3F_7 group;
and r represents an integer selected within a range

- 195 -

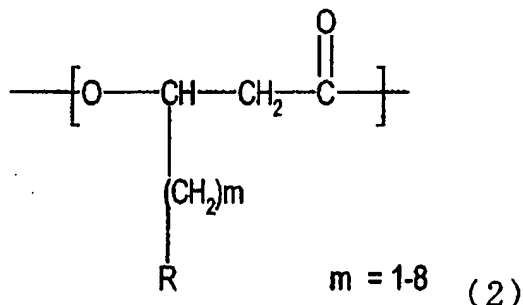
indicated in the chemical formula;

[Chemical Formula (1)]



in which n represents an integer selected within a
 5 range indicated in the chemical formula; and in case
 plural units are present, n is the same or different
 for each unit;

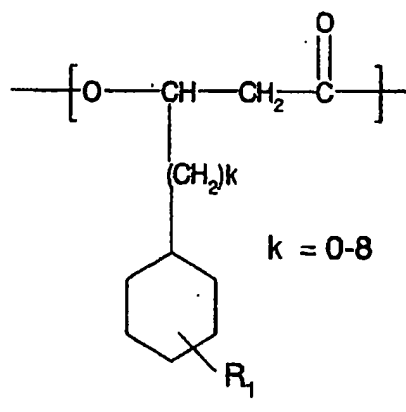
[Chemical Formula (2)]



10 in which m represents an integer selected within a
 range indicated in the chemical formula; R represents
 a residue having any of a phenyl structure or a
 thienyl structure; and in case plural units are
 present, m and R are the same or different for each
 15 unit; and

- 196 -

[Chemical Formula (3)]

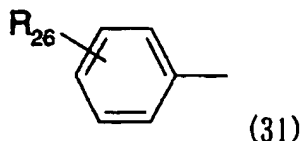


in which R_1 represents a substituent on a cyclohexyl group and represents an H atom, a CN group, a NO_2 group, a halogen atom, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a CF_3 group, a C_2F_5 group, or a C_3F_7 group; k represents an integer selected within a range indicated in the chemical formula; and in case plural units are present, R_1 and k are the same or different for each unit.

8. The method for producing a polyhydroxy alkanoate copolymer according to claim 7, wherein R_{23} in the chemical formula (25) and R in the chemical formula (2), each represents a residue having a phenyl structure or a thienyl structure, are selected from chemical formulas (31), (9), (10), (11), (12), (13), (14), (15), (16), (17) and (18):

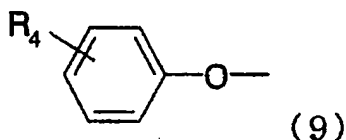
the chemical formula (31):

- 197 -



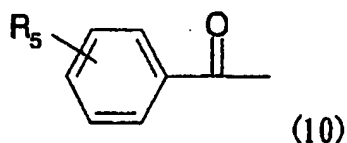
represents a group of substituted or non-substituted phenyl groups in which R_{26} represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $CH=CH_2$ group, a CF_3 group, a C_2F_5 group or a C_3F_7 group; and in case plural units are present, R_{26} is the same or different for each unit;

the chemical formula (9):



represents a group of non-substituted or substituted phenoxy groups in which R_4 represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a SCH_3 group, a CF_3 group, a C_2F_5 group, or a C_3F_7 group; and in case plural units are present, R_4 is the same or different for each unit;

the chemical formula (10):

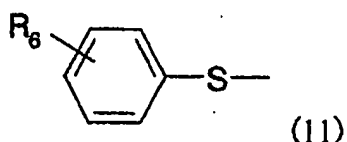


represents a group of non-substituted or substituted

- 198 -

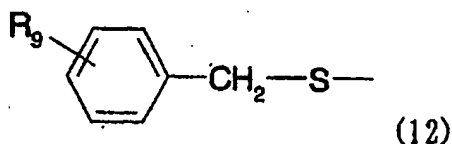
benzoyl groups in which R_5 represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a CF_3 group, a C_2F_5 group, or a
 5 C_3F_7 group; and in case plural units are present, R_5 is the same or different for each unit;

the chemical formula (11):



represents a group of substituted or non-substituted
 10 phenylsulfanyl groups in which R_6 represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a COOR_7 group, a SO_2R_8 group (R_7 representing either one of H, Na, K, CH_3 and C_2H_5 ; and R_8 representing either one of
 15 OH, ONa, OK, a halogen atom, OCH_3 and OC_2H_5), a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $(\text{CH}_3)_2\text{-CH}$ group or a $(\text{CH}_3)_3\text{-C}$ group; and in case plural units are present, R_6 is the same or different for each unit;

the chemical formula (12):



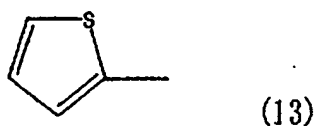
20

represents a group of substituted or non-substituted (phenylmethyl)sulfanyl groups in which R_9 represents a substituent on an aromatic ring and represents an H

- 199 -

atom, a halogen atom, a CN group, a NO₂ group, a COOR₁₀ group, a SO₂R₁₁ group (R₁₀ representing either one of H, Na, K, CH₃ and C₂H₅; and R₁₁ representing either one of OH, ONa, OK, a halogen atom, OCH₃ and OC₂H₅), a CH₃ group, a C₂H₅ group, a C₃H₇ group, a (CH₃)₂-CH group or a (CH₃)₃-C group; and in case plural units are present, R₉ is the same or different for each unit;

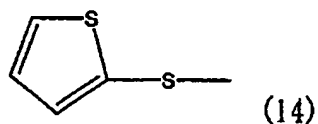
the chemical formula (13):



10

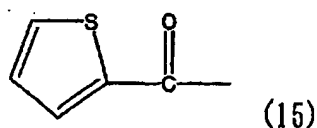
represents a 2-thienyl group;

the chemical formula (14):



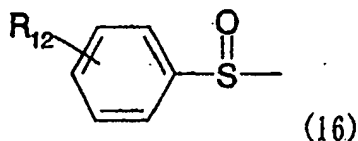
represents a 2-thienylsulfanyl group;

15 the chemical formula (15):



represents a 2-thienylcarbonyl group;

the chemical formula (16):



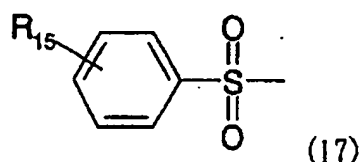
- 200 -

represents a group of substituted or non-substituted phenylsulfinyl groups in which R_{12} represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a

5 COOR_{13} group, a SO_2R_{14} group (R_{13} representing either one of H, Na, K, CH_3 and C_2H_5 ; and R_{14} representing either one of OH, ONa, OK, a halogen atom, OCH_3 and OC_2H_5), a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $(\text{CH}_3)_2\text{-CH}$ group or a $(\text{CH}_3)_3\text{-C}$ group; and in case

10 plural units are present, R_{12} is the same or different for each unit;

the chemical formula (17):



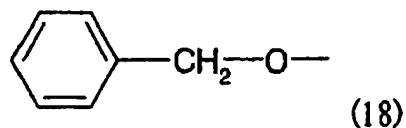
represents a group of substituted or non-substituted

15 phenylsulfonyl groups in which R_{15} represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a COOR_{16} group, a SO_2R_{17} group (R_{16} representing either one of H, Na, K, CH_3 and C_2H_5 ; and R_{17} representing

20 either one of OH, ONa, OK, a halogen atom, OCH_3 and OC_2H_5), a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $(\text{CH}_3)_2\text{-CH}$ group or a $(\text{CH}_3)_3\text{-C}$ group; and in case plural units are present, R_{15} is the same or different for each unit; and

25 the chemical formula (18):

- 201 -



represents a (phenylmethyl)oxy group.

9. The method for producing a polyhydroxy
5 alkanolate copolymer according to claim 7, wherein
said microorganism is cultured in a culture medium
including at least an ω -alkenoic acid represented by
the chemical formula (24) and at least a compound
represented by the chemical formula (25) or at least
10 an ω -cyclohexylalkanoic acid represented by the
chemical formula (26).

10. The method for producing a polyhydroxy
alkanoate copolymer according to claim 9, wherein
15 said microorganism is cultured in a culture medium
including, in addition to at least an ω -alkenoic acid
represented by the chemical formula (24) and at least
a compound represented by the chemical formula (25)
or at least an ω -cyclohexylalkanoic acid represented
20 by the chemical formula (26), at least one of a
peptide, an yeast extract, an organic acid or a salt
thereof, an amino acid or a salt thereof, a sugar, a
linear alkanolic acid with 4 to 12 carbon atoms or a
salt thereof.

- 202 -

11. The method for producing a polyhydroxy
alkanoate copolymer according to claim 7,
characterized in including a step of culturing said
microorganism in a culture medium including at least
5 an ω -alkenoic acid represented by the chemical
formula (24) and at least a compound represented by
the chemical formula (25) or at least an ω -
cyclohexylalkanoic acid represented by the chemical
formula (26), and recovering a polyhydroxy alkanoate
10 copolymer including simultaneously at least a 3-
hydroxy- ω -alkenoic acid unit represented by the
chemical formula (1) and a 3-hydroxy- ω -alkanoic acid
unit represented by the chemical formula (2) or a 3-
hydroxy- ω -cyclohexylalkanoic acid unit represented by
15 the chemical formula (3) in the molecule, produced by
said microorganism, from cells of the microorganism.

12. The method for producing a polyhydroxy
alkanoate copolymer according to claim 7, wherein
20 said microorganism is a microorganism belonging to
Pseudomonas genus.

13. The method for producing a polyhydroxy
alkanoate copolymer according to claim 12, wherein
25 said microorganism is at least one of *Pseudomonas*
cichorii YN2 strain (FERM BP-7375), *Pseudomonas*
cichorii H45 strain (FERM BP-7374), *Pseudomonas*

- 203 -

jessenii P161 (FERM BP-7376) and *Pseudomonas putida* P91 (FERM BP-7373).

14. A method for producing a polyhydroxy
5 alkanooate copolymer including at least a 3-hydroxy- ω -
carboxyalkanoic acid unit represented by a chemical
formula (19) in a molecule, and simultaneously at
least a 3-hydroxy- ω -alkanoic acid unit represented by
a chemical formula (2) or a 3-hydroxy- ω -
10 cyclohexylalkanoic acid unit represented by a
chemical formula (3) in the molecule comprising the
steps of:

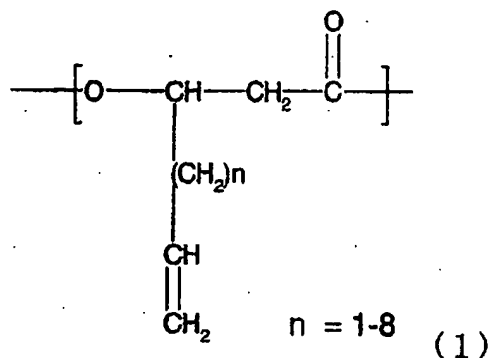
preparing a polyhydroxy alkanooate copolymer
including at least a 3-hydroxy- ω -alkenoic acid unit
15 represented by a chemical formula (1) in a molecule,
and simultaneously at least a 3-hydroxy- ω -alkanoic
acid unit represented by a chemical formula (2) or a
3-hydroxy- ω -cyclohexylalkanoic acid unit represented
by a chemical formula (3) in the molecule as a
20 starting material, and

oxidizing a double bond portion in the
polyhydroxy alkanooate represented in the chemical
formula (1) thereby generating a polyhydroxy
alkanoate copolymer including at least a 3-hydroxy- ω -
25 carboxyalkanoic acid unit represented by a chemical
formula (19) in a molecule, and simultaneously at
least a 3-hydroxy- ω -alkanoic acid unit represented by

- 204 -

a chemical formula (2) or a 3-hydroxy- ω -cyclohexylalkanoic acid unit represented by a chemical formula (3) in the molecule:

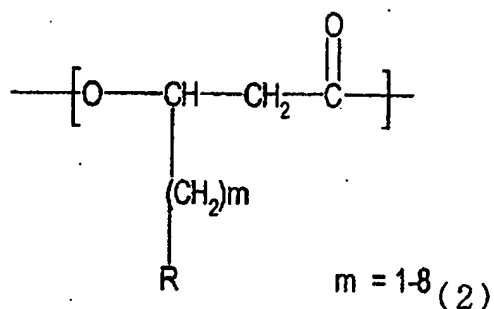
[Chemical Formula (1)]



5

in which n represents an integer selected within a range indicated in the chemical formula; and in case plural units are present, n is the same or different for each unit;

10 [Chemical Formula (2)]

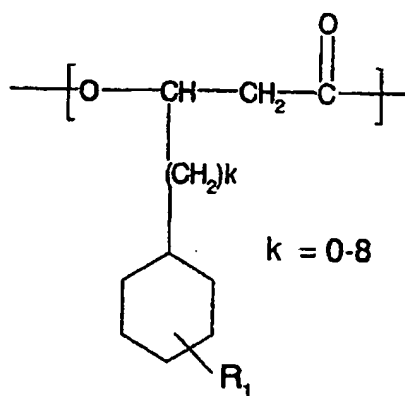


in which m represents an integer selected within a range indicated in the chemical formula; R includes a residue having any of a phenyl structure and a
 15 thienyl structure; and in case plural units are present, m and R are the same or different for each

- 205 -

unit;

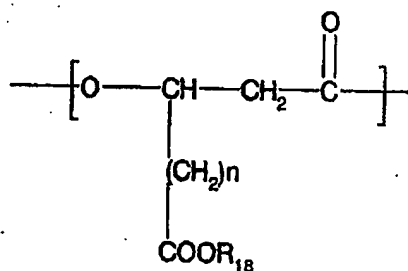
[Chemical Formula (3)]



(3)

in which R_1 represents a substituent on a cyclohexyl
 5 group selected from an H atom, a CN group, a NO_2
 group, a halogen atom, a CH_3 group, a C_2H_5 group, a
 C_3H_7 group, a CF_3 group, a C_2F_5 group, and a C_3F_7
 group; k represents an integer selected within a
 range indicated in the chemical formula; and in case
 10 plural units are present, R_1 and k are the same or
 different for each unit; and

[Chemical Formula (19)]

 $n = 1-8$ (19)

in which n represents an integer selected within a

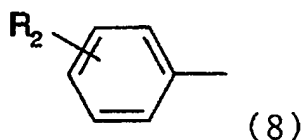
- 206 -

range indicated in the chemical formula; R_{18} represents an H atom, a Na atom, or a K atom; and in case plural units are present, n and R_{18} are the same or different for each unit.

5

15. The method for producing a polyhydroxy alkanoate copolymer according to claim 14, wherein R in the chemical formula (2) represents a residue having a phenyl structure or a thienyl structure
10 selected from chemical formulas (8), (9), (10), (11), (12), (13), (14), (15), (16), (17) and (18):

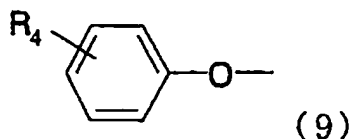
the chemical formula (8):



represents a group of non-substituted or substituted
15 phenyl groups in which R_2 represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $\text{CH}=\text{CH}_2$ group, a COOR_3 group (R_3 representing an H atom, a Na atom or a K atom), a CF_3
20 group, a C_2F_5 group, or a C_3F_7 group; and in case plural units are present, R_2 is the same or different for each unit;

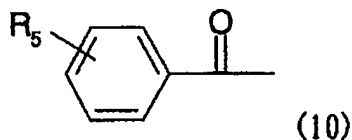
the chemical formula (9):

- 207 -



represents a group of non-substituted or substituted phenoxy groups in which R_4 represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a SCH_3 group, a CF_3 group, a C_2F_5 group, or a C_3F_7 group; and in case plural units are present, R_4 is the same or different for each unit;

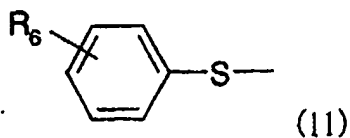
the chemical formula (10):



10

represents a group of non-substituted or substituted benzoyl groups in which R_5 represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a CF_3 group, a C_2F_5 group, or a C_3F_7 group; and in case plural units are present, R_5 is the same or different for each unit;

the chemical formula (11):

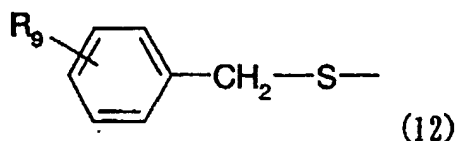


- 208 -

represents a group of substituted or non-substituted phenylsulfanyl groups in which R_6 represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a $COOR_7$ group, a SO_2R_8 group (R_7 represents either one of H, Na, K, CH_3 and C_2H_5 ; and R_8 represents either one of OH, ONa, OK, a halogen atom, OCH_3 and OC_2H_5), a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $(CH_3)_2-CH$ group or a $(CH_3)_3-C$ group; and in case plural units are present,

10 R_6 is the same or different for each unit;

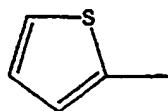
the chemical formula (12):



represents a group of substituted or non-substituted (phenylmethyl)sulfanyl groups in which R_9 represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a $COOR_{10}$ group, a SO_2R_{11} group (R_{10} represents either one of H, Na, K, CH_3 and C_2H_5 ; and R_{11} represents either one of OH, ONa, OK, a halogen atom, OCH_3 and OC_2H_5), a

15 CH_3 group, a C_2H_5 group, a C_3H_7 group, a $(CH_3)_2-CH$ group or a $(CH_3)_3-C$ group; and in case plural units are present, R_9 is the same or different for each unit;

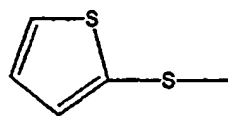
the chemical formula (13):



(13)

represents a 2-thienyl group;

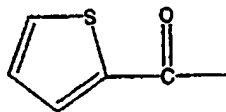
the chemical formula (14)



(14)

5 represents a 2-thienylsulfanyl group;

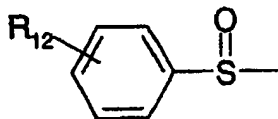
the chemical formula (15):



(15)

represents a 2-thienylcarbonyl group;

the chemical formula (16):



(16)

10

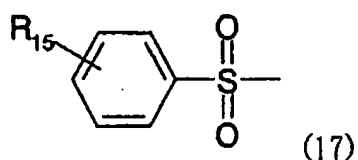
represents a group of substituted or non-substituted phenylsulfinyl groups in which R_{12} represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a

15 $COOR_{13}$ group, a SO_2R_{14} group (R_{13} represents either one of H, Na, K, CH_3 and C_2H_5 ; and R_{14} represents either one of OH, ONa, OK, a halogen atom, OCH_3 and OC_2H_5), a

- 210 -

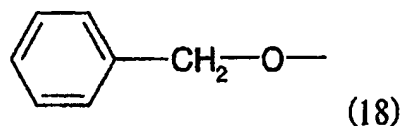
CH₃ group, a C₂H₅ group, a C₃H₇ group, a (CH₃)₂-CH group or a (CH₃)₃-C group; and in case plural units are present, R₁₂ is the same or different for each unit;

5 the chemical formula (17):



represents a group of substituted or non-substituted phenylsulfonyl groups in which R₁₅ represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO₂ group, a COOR₁₆ group, a SO₂R₁₇ group (R₁₆ represents either one of H, Na, K, CH₃ and C₂H₅; and R₁₇ represents either one of OH, ONa, OK, a halogen atom, OCH₃ and OC₂H₅), a CH₃ group, a C₂H₅ group, a C₃H₇ group, a (CH₃)₂-CH group or a (CH₃)₃-C group; and in case plural units are present, R₁₅ is the same or different for each unit;

the chemical formula (18):



20 represents a (phenylmethyl)oxy group.

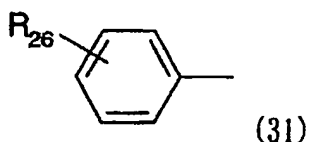
16. The method according to claim 14, wherein said starting material polyhydroxy alkanoate

- 211 -

copolymer including at least a 3-hydroxy- ω -alkenoic acid unit represented by a chemical formula (1) in a molecule, and simultaneously at least a 3-hydroxy- ω -alkanoic acid unit represented by a chemical formula
 5 (2) or a 3-hydroxy- ω -cyclohexylalkanoic acid unit represented by a chemical formula (3) in the molecule, is produced by a method according to claim 7.

17. The method for producing a polyhydroxy
 10 alkanoate copolymer according to claim 16, wherein R in the chemical formula (2), representing a residue having a phenyl structure or a thienyl structure, is at least one of chemical formulas (31), (9), (10), (11), (12), (13), (14), (15), (16), (17) and (18):

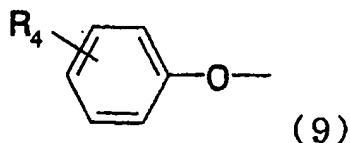
15 the chemical formula (31):



represents a group of substituted or non-substituted phenyl groups in which R_{26} represents a substituent on an aromatic ring and represents an H atom, a halogen
 20 atom, a CN group, a NO_2 group, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $CH=CH_2$ group, a CF_3 group, a C_2F_5 group or a C_3F_7 group; and in case plural units are present, R_{26} is the same or different for each unit;

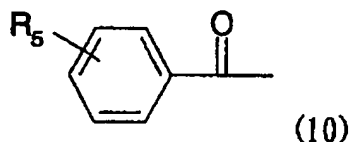
25 the chemical formula (9):

- 212 -



represents a group of non-substituted or substituted phenoxy groups in which R_4 represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a SCH_3 group, a CF_3 group, a C_2F_5 group, or a C_3F_7 group; and in case plural units are present, R_4 is the same or different for each unit;

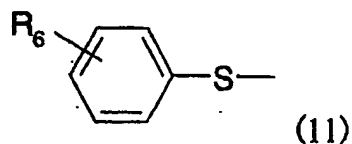
the chemical formula (10):



10

represents a group of non-substituted or substituted benzoyl groups in which R_5 represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a CF_3 group, a C_2F_5 group, or a C_3F_7 group; and in case plural units are present, R_5 is the same or different for each unit;

the chemical formula (11):

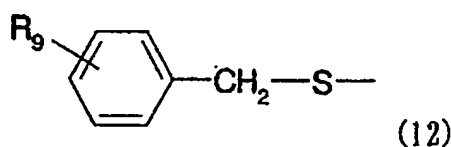


20 represents a group of substituted or non-substituted phenylsulfanyl groups in which R_6 represents a

- 213 -

substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO₂ group, a COOR₇ group, a SO₂R₈ group (R₇ representing either one of H, Na, K, CH₃ and C₂H₅; and R₈ representing either one of
 5 OH, ONa, OK, a halogen atom, OCH₃ and OC₂H₅), a CH₃ group, a C₂H₅ group, a C₃H₇ group, a (CH₃)₂-CH group or a (CH₃)₃-C group; and in case plural units are present, R₆ is the same or different for each unit;

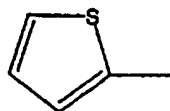
the chemical formula (12):



10

represents a group of substituted or non-substituted (phenylmethyl)sulfanyl groups in which R₉ represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO₂ group, a
 15 COOR₁₀ group, a SO₂R₁₁ group (R₁₀ representing either one of H, Na, K, CH₃ and C₂H₅; and R₁₁ representing either one of OH, ONa, OK, a halogen atom, OCH₃ and OC₂H₅), a CH₃ group, a C₂H₅ group, a C₃H₇ group, a (CH₃)₂-CH group or a (CH₃)₃-C group; and in case
 20 plural units are present, R₉ is the same or different for each unit;

the chemical formula (13):

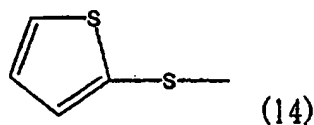


(13)

- 214 -

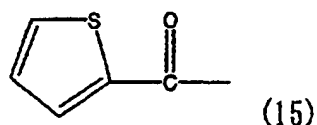
represents a 2-thienyl group;

the chemical formula (14):



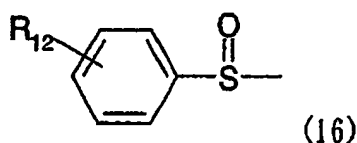
represents a 2-thienylsulfanyl group;

5 the chemical formula (15):



represents a 2-thienylcarbonyl group;

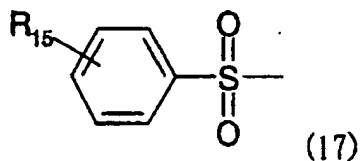
the chemical formula (16):



10 represents a group of substituted or non-substituted phenylsulfanyl groups in which R_{12} represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a $COOR_{13}$ group, a SO_2R_{14} group (R_{13} representing either
 15 one of H, Na, K, CH_3 and C_2H_5 ; and R_{14} representing either one of OH, ONa, OK, a halogen atom, OCH_3 and OC_2H_5), a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $(CH_3)_2-CH$ group or a $(CH_3)_3-C$ group; and in case plural units are present, R_{12} is the same or different
 20 for each unit;

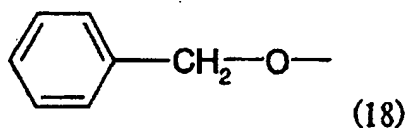
the chemical formula (17):

- 215 -



represents a group of substituted or non-substituted phenylsulfonyl groups in which R_{15} represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a $COOR_{16}$ group, a SO_2R_{17} group (R_{16} representing either one of H, Na, K, CH_3 and C_2H_5 ; and R_{17} representing either one of OH, ONa, OK, a halogen atom, OCH_3 and OC_2H_5), a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $(CH_3)_2-CH$ group or a $(CH_3)_3-C$ group; and in case plural units are present, R_{15} is the same or different for each unit; and

the chemical formula (18):



represents a (phenylmethyl)oxy group.

18. The producing method according to claim 14, wherein said oxidation reaction is carried out with an oxidant selected from a group consisting of a permanganate, a bichromate and a periodate.

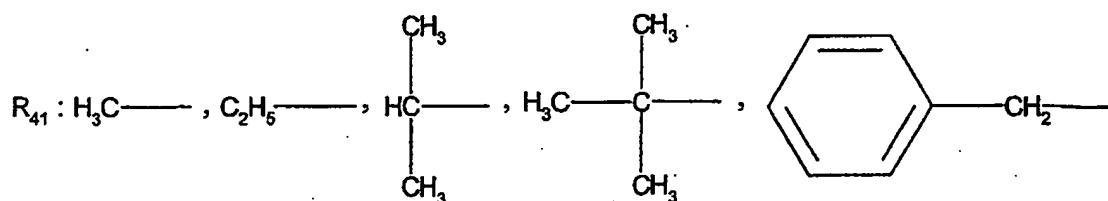
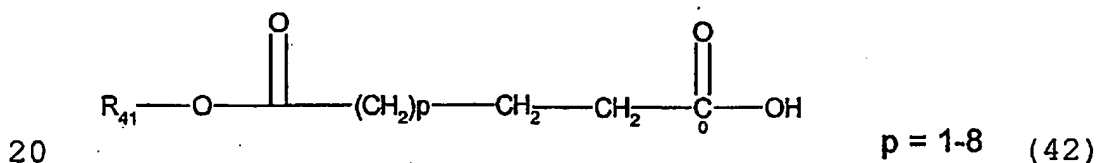
19. The producing method according to claim 18, wherein said oxidation reaction is carried out with a

- 216 -

permanganate as an oxidant and under an acidic condition.

20. The producing method according to claim 14,
5 wherein said oxidation reaction is carried out with ozone.

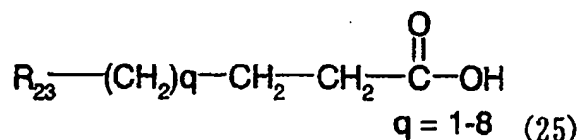
21. The method for producing a polyhydroxy
alkanoate copolymer including a biosynthesis by a
10 microorganism having an ability of producing a
polyhydroxy alkanoate copolymer including at least a
3-hydroxy- ω -alkoxycarbonylalkanoic acid unit
represented by a chemical formula (32) in a molecule,
and simultaneously at least a 3-hydroxy- ω -alkanoic
15 acid unit represented by a chemical formula (2) or a
3-hydroxy- ω -cyclohexylalkanoic acid unit represented
by a chemical formula (3) in the molecule, from a
dicarboxylic acid monoester compound represented by a
chemical formula (42):



- 217 -

in which p may assume one or more arbitrary integral values within a range indicated in the chemical formula; and R₄₁ may arbitrarily represent one or more residues indicated in the chemical formula; and at least a compound represented by a chemical formula (25) or at least a ω-cyclohexylalkanoic acid represented by a chemical formula (26) as starting materials:

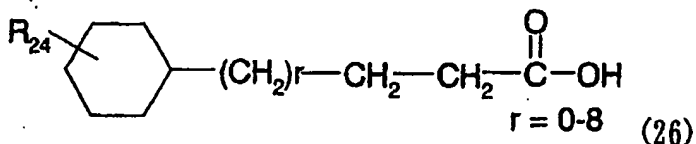
[Chemical Formula (25)]



10

in which q represents an integer selected within a range indicated in the chemical formula; and R₂₃ includes a residue having a phenyl structure or a thienyl structure;

15 [Chemical Formula (26)]



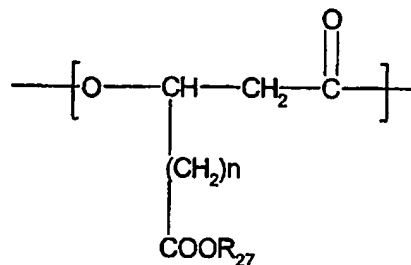
in which R₂₄ represents a substituent on a cyclohexyl group and represents an H atom, a CN group, a NO₂ group, a halogen atom, a CH₃ group, a C₂H₅ group, a C₃H₇ group, a CF₃ group, a C₂F₅ group, or a C₃F₇ group; and r represents an integer selected within a range

20

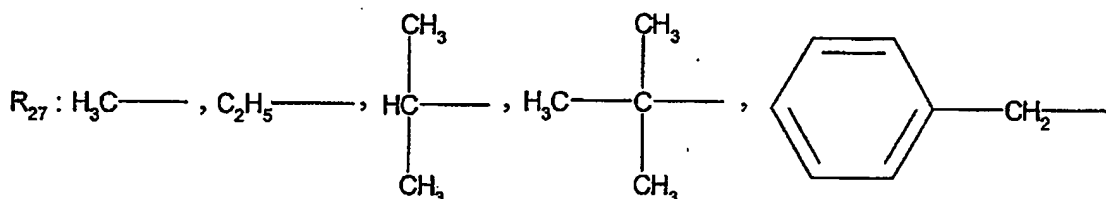
- 218 -

indicated in the chemical formula;

[Chemical Formula (32)]



$n = 1-8$ (32)



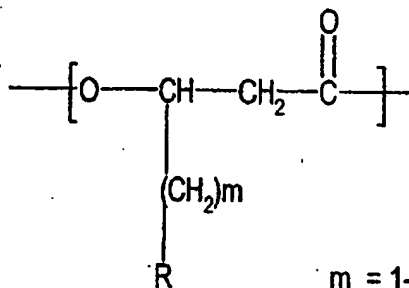
5

in which n represents an integer selected within a range indicated in the chemical formula; R_{27}

represents any of residues indicated in the chemical formula; and in case plural units are present, n and

10 R_{27} are the same or different for each unit;

[Chemical Formula (2)]



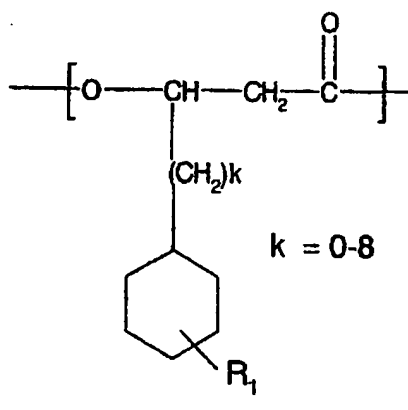
$m = 1-8$ (2)

in which m represents an integer selected within a range indicated in the chemical formula; R represents

- 219 -

a residue having any of a phenyl structure and a thienyl structure; and in case plural units are present, m and R are the same or different for each unit; and

5 [Chemical Formula (3)]



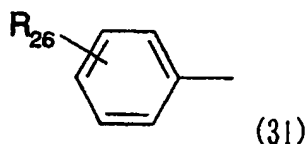
in which R_1 represents a substituent on a cyclohexyl group and represents an H atom, a CN group, a NO_2 group, a halogen atom, a CH_3 group, a C_2H_5 group, a
 10 C_3H_7 group, a CF_3 group, a C_2F_5 group, or a C_3F_7 group;
 k represents an integer selected within a range indicated in the chemical formula; and in case plural units are present, R_1 and k are the same or different for each unit.

15

22. The method for producing a polyhydroxy alkanooate copolymer according to claim 21, wherein R_{23} in the chemical formula (25) and R in the chemical formula (2), each representing a residue having a
 20 phenyl structure or a thienyl structure, represents at least one of chemical formulas (31), (9), (10),

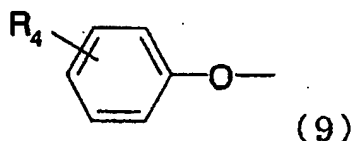
- 220 -

(11), (12), (13), (14), (15), (16), (17) and (18):
the chemical formula (31):



represents a group of substituted or non-substituted
5 phenyl groups in which R_{26} represents a substituent on
an aromatic ring and represents an H atom, a halogen
atom, a CN group, a NO_2 group, a CH_3 group, a C_2H_5
group, a C_3H_7 group, a $\text{CH}=\text{CH}_2$ group, a CF_3 group, a
 C_2F_5 group or a C_3F_7 group; and in case plural units
10 are present, R_{26} is the same or different for each
unit;

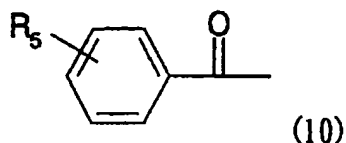
the chemical formula (9):



represents a group of non-substituted or substituted
15 phenoxy groups in which R_4 represents a substituent on
an aromatic ring and represents an H atom, a halogen
atom, a CN group, a NO_2 group, a CH_3 group, a C_2H_5
group, a C_3H_7 group, a SCH_3 group, a CF_3 group, a C_2F_5
group, or a C_3F_7 group; and in case plural units are
20 present, R_4 is the same or different for each unit;

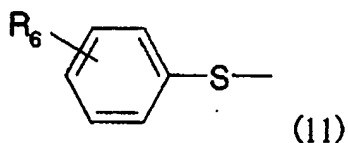
the chemical formula (10):

- 221 -



represents a group of non-substituted or substituted benzoyl groups in which R_5 represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a CF_3 group, a C_2F_5 group, or a C_3F_7 group; and in case plural units are present, R_5 is the same or different for each unit;

the chemical formula (11):



10

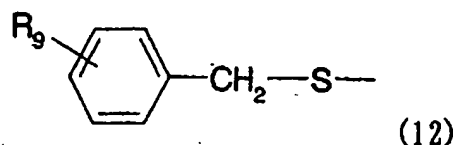
represents a group of substituted or non-substituted phenylsulfanyl groups in which R_6 represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a $COOR_7$ group, a SO_2R_8 group (R_7 representing either one of H, Na, K, CH_3 and C_2H_5 ; and R_8 representing either one of OH, ONa, OK, a halogen atom, OCH_3 and OC_2H_5), a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $(CH_3)_2-CH$ group or a $(CH_3)_3-C$ group; and in case plural units are present,

20

R_6 is the same or different for each unit;

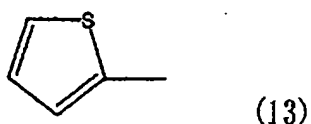
the chemical formula (12):

- 222 -



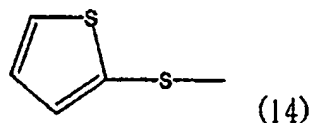
represents a group of substituted or non-substituted (phenylmethyl)sulfanyl groups in which R_9 represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a COOR_{10} group, a SO_2R_{11} group (R_{10} representing either one of H, Na, K, CH_3 and C_2H_5 ; and R_{11} representing either one of OH, ONa, OK, a halogen atom, OCH_3 and OC_2H_5), a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $(\text{CH}_3)_2\text{-CH}$ group or a $(\text{CH}_3)_3\text{-C}$ group; and in case plural units are present, R_9 is the same or different for each unit;

the chemical formula (13):



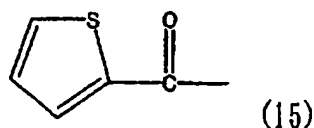
15 represents a 2-thienyl group;

the chemical formula (14):



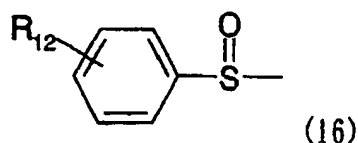
represents a 2-thienylsulfanyl group;

the chemical formula (15):



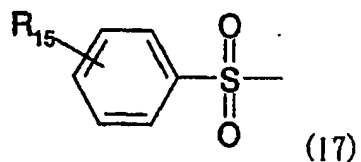
represents a 2-thienylcarbonyl group;

the chemical formula (16):



5 represents a group of substituted or non-substituted phenylsulfinyl groups in which R_{12} represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a COOR_{13} group, a SO_2R_{14} group (R_{13} representing either
 10 one of H, Na, K, CH_3 and C_2H_5 ; and R_{14} representing either one of OH, ONa, OK, a halogen atom, OCH_3 and OC_2H_5), a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $(\text{CH}_3)_2\text{-CH}$ group or a $(\text{CH}_3)_3\text{-C}$ group; and in case plural units are present, R_{12} is the same or different
 15 for each unit;

the chemical formula (17):

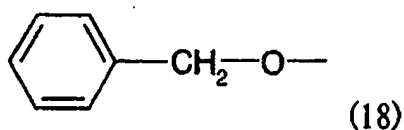


represents a group of substituted or non-substituted phenylsulfonyl groups in which R_{15} represents a
 20 substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a

- 224 -

COOR₁₆ group, a SO₂R₁₇ group (R₁₆ representing either one of H, Na, K, CH₃ and C₂H₅; and R₁₇ representing either one of OH, ONa, OK, a halogen atom, OCH₃ and OC₂H₅), a CH₃ group, a C₂H₅ group, a C₃H₇ group, a
 5 (CH₃)₂-CH group or a (CH₃)₃-C group; and in case plural units are present, R₁₅ is the same or different for each unit; and

the chemical formula (18):



10 represents a (phenylmethyl)oxy group.

23. The method for producing a polyhydroxy alkanate copolymer according to claim 21, wherein the microorganism is cultured in a culture medium
 15 including at least a dicarboxylic acid monoester compound represented by the chemical formula (42) and at least a compound represented by the chemical formula (25) or at least an ω-cyclohexylalkanoic acid represented by the chemical formula (26).

20

24. The method for producing a polyhydroxy alkanate copolymer according to claim 23, wherein the microorganism is cultured in a culture medium including, in addition, at least one of a peptide, an
 25 yeast extract, an organic acid or a salt thereof, an

- 225 -

amino acid or a salt thereof, a sugar, a linear alkanolic acid with 4 to 12 carbon atoms or a salt thereof.

5 25. The method for producing a polyhydroxy alkanolate copolymer according to claim 21, characterized in including a step of recovering a polyhydroxy alkanolate copolymer, produced by said microorganism, from cells of the microorganism.

10

26. The method for producing a polyhydroxy alkanolate copolymer according to claim 21, wherein said microorganism is a microorganism belonging to *Pseudomonas* genus.

15

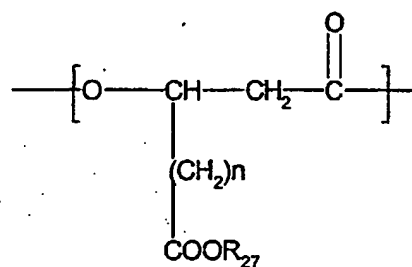
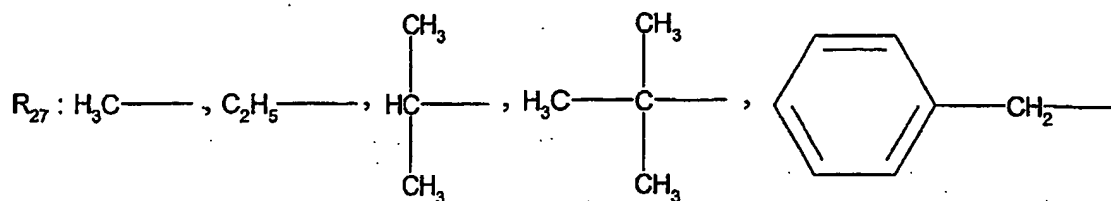
27. The method for producing a polyhydroxy alkanolate copolymer according to claim 26, wherein said microorganism is at least one of *Pseudomonas cichorii* YN2 strain (FERM BP-7375), *Pseudomonas*
20 *cichorii* H45 strain (FERM BP-7374), *Pseudomonas jessenii* P161 (FERM BP-7376) and *Pseudomonas putida* P91 (FERM BP-7373).

28. A method for producing a polyhydroxy
25 alkanolate copolymer, characterized in employing a polyhydroxy alkanolate copolymer including at least a 3-hydroxy- ω -alkoxycarbonylalkanoic acid unit

- 226 -

represented by a chemical formula (32) in a molecule,
 and simultaneously at least a 3-hydroxy- ω -alkanoic
 acid unit represented by a chemical formula (2) or a
 3-hydroxy- ω -cyclohexylalkanoic acid unit represented
 5 by a chemical formula (3) in the molecule as a
 starting material, and executing a hydrolysis in the
 presence of an acid or an alkali or executing a
 hydrogenolysis including a catalytic reduction,
 thereby generating a polyhydroxy alkanoate copolymer
 10 including at least a 3-hydroxy- ω -carboxyalkanoic acid
 unit represented by a chemical formula (19) in a
 molecule, and simultaneously at least a 3-hydroxy- ω -
 alkanoic acid unit represented by a chemical formula
 (2) or a 3-hydroxy- ω -cyclohexylalkanoic acid unit
 15 represented by a chemical formula (3) in the
 molecule:

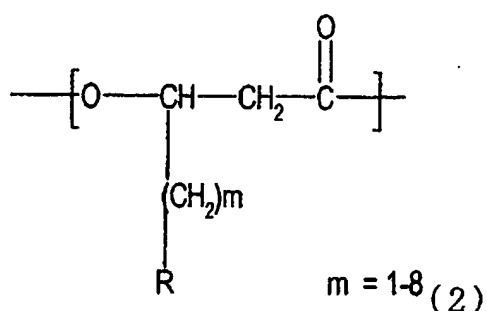
[Chemical Formula (32)]

 $n = 1-8$ (32)

- 227 -

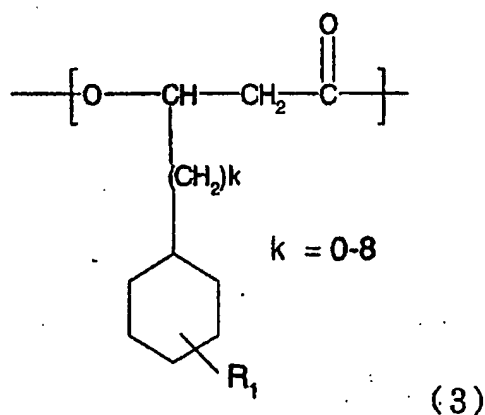
in which n represents an integer selected within a range indicated in the chemical formula; R_{27} represents any of residues indicated in the chemical formula; and in case plural units are present, n and R_{27} are the same or different for each unit;

[Chemical Formula (2)]



in which m represents an integer selected within a range indicated in the chemical formula; R includes a residue having any of a phenyl structure and a thienyl structure; and in case plural units are present, m and R are the same or different for each unit;

[Chemical Formula (3)]

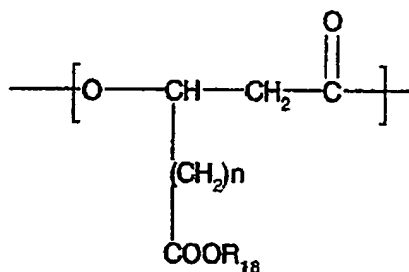


- 228 -

in which R_1 represents a substituent on a cyclohexyl group and represents an H atom, a CN group, a NO_2 group, a halogen atom, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a CF_3 group, a C_2F_5 group, or a C_3F_7 group;

5 k represents an integer selected within a range indicated in the chemical formula; and in case plural units are present, R_1 and k are the same or different for each unit; and

[Chemical Formula (19)]



$$n = 1-8 \quad (19)$$

10

in which n represents an integer selected within a range indicated in the chemical formula; R_{18} represents an H atom, a Na atom, or a K atom; and in case plural units are present, n and R_{18} are the same

15 or different for each unit.

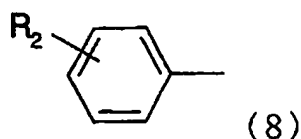
29. The method for producing a polyhydroxy alkanooate copolymer according to claim 28, wherein R in the chemical formula (2), representing a residue

20 having a phenyl structure or a thienyl structure, represents at least one of chemical formulas (8), (9),

- 229 -

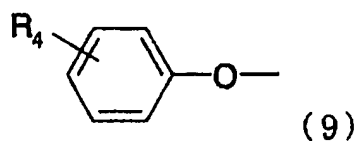
(10), (11), (12), (13), (14), (15), (16), (17) and (18):

the chemical formula (8):



5 represents a group of non-substituted or substituted phenyl groups in which R_2 represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $\text{CH}=\text{CH}_2$ group, a COOR_3 group (R_3 representing an H atom, a Na atom or a K atom), a CF_3 group, a C_2F_5 group, or a C_3F_7 group; and in case plural units are present, R_2 is the same or different for each unit;

the chemical formula (9):



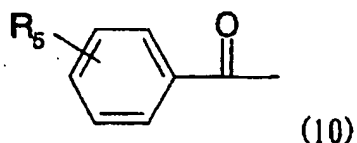
15

represents a group of non-substituted or substituted phenoxy groups in which R_4 represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a SCH_3 group, a CF_3 group, a C_2F_5 group, or a C_3F_7 group; and in case plural units are

20

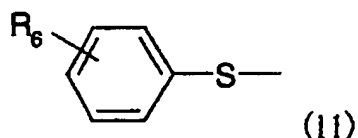
- 230 -

present, R_4 is the same or different for each unit;
the chemical formula (10):



represents a group of non-substituted or substituted
5 benzoyl groups in which R_5 represents a substituent on
an aromatic ring and represents an H atom, a halogen
atom, a CN group, a NO_2 group, a CH_3 group, a C_2H_5
group, a C_3H_7 group, a CF_3 group, a C_2F_5 group, or a
 C_3F_7 group; and in case plural units are present, R_5
10 is the same or different for each unit;

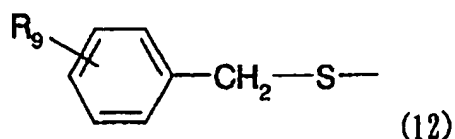
the chemical formula (11):



represents a group of substituted or non-substituted
phenylsulfanyl groups in which R_6 represents a
15 substituent on an aromatic ring and represents an H
atom, a halogen atom, a CN group, a NO_2 group, a COOR_7
group, a SO_2R_8 group (R_7 represents either one of H,
Na, K, CH_3 and C_2H_5 ; and R_8 represents either one of
OH, ONa, OK, a halogen atom, OCH_3 and OC_2H_5), a CH_3
20 group, a C_2H_5 group, a C_3H_7 group, a $(\text{CH}_3)_2\text{-CH}$ group or
a $(\text{CH}_3)_3\text{-C}$ group; and in case plural units are present,

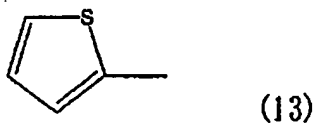
- 231 -

R_6 is the same or different for each unit;
the chemical formula (12):



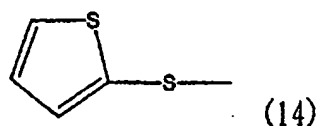
represents a group of substituted or non-substituted
5 (phenylmethyl)sulfanyl groups in which R_9 represents a
substituent on an aromatic ring and represents an H
atom, a halogen atom, a CN group, a NO_2 group, a
 COOR_{10} group, a SO_2R_{11} group (R_{10} represents either one
of H, Na, K, CH_3 and C_2H_5 ; and R_{11} represents either
10 one of OH, ONa, OK, a halogen atom, OCH_3 and OC_2H_5), a
 CH_3 group, a C_2H_5 group, a C_3H_7 group, a $(\text{CH}_3)_2\text{-CH}$
group or a $(\text{CH}_3)_3\text{-C}$ group; and in case plural units
are present, R_9 is the same or different for each
unit;

15 the chemical formula (13):



represents a 2-thienyl group;

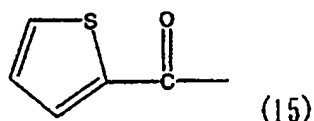
the chemical formula (14):



- 232 -

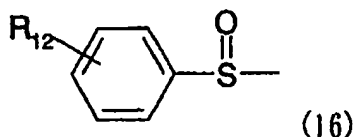
represents a 2-thienylsulfanyl group;

the chemical formula (15):



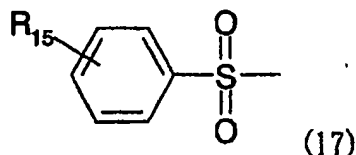
represents a 2-thienylcarbonyl group;

5 the chemical formula (16):



represents a group of substituted or non-substituted phenylsulfinyl groups in which R_{12} represents a substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO_2 group, a COOR_{13} group, a SO_2R_{14} group (R_{13} represents either one of H, Na, K, CH_3 and C_2H_5 ; and R_{14} represents either one of OH, ONa, OK, a halogen atom, OCH_3 and OC_2H_5), a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $(\text{CH}_3)_2\text{-CH}$ group or a $(\text{CH}_3)_3\text{-C}$ group; and in case plural units are present, R_{12} is the same or different for each unit;

the chemical formula (17):

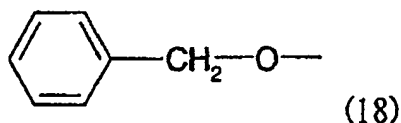


20 represents a group of substituted or non-substituted phenylsulfonyl groups in which R_{15} represents a

- 233 -

substituent on an aromatic ring and represents an H atom, a halogen atom, a CN group, a NO₂ group, a COOR₁₆ group, a SO₂R₁₇ group (R₁₆ represents either one of H, Na, K, CH₃ and C₂H₅; and R₁₇ represents either
5 one of OH, ONa, OK, a halogen atom, OCH₃ and OC₂H₅), a CH₃ group, a C₂H₅ group, a C₃H₇ group, a (CH₃)₂-CH group or a (CH₃)₃-C group; and in case plural units are present, R₁₅ is the same or different for each unit; and

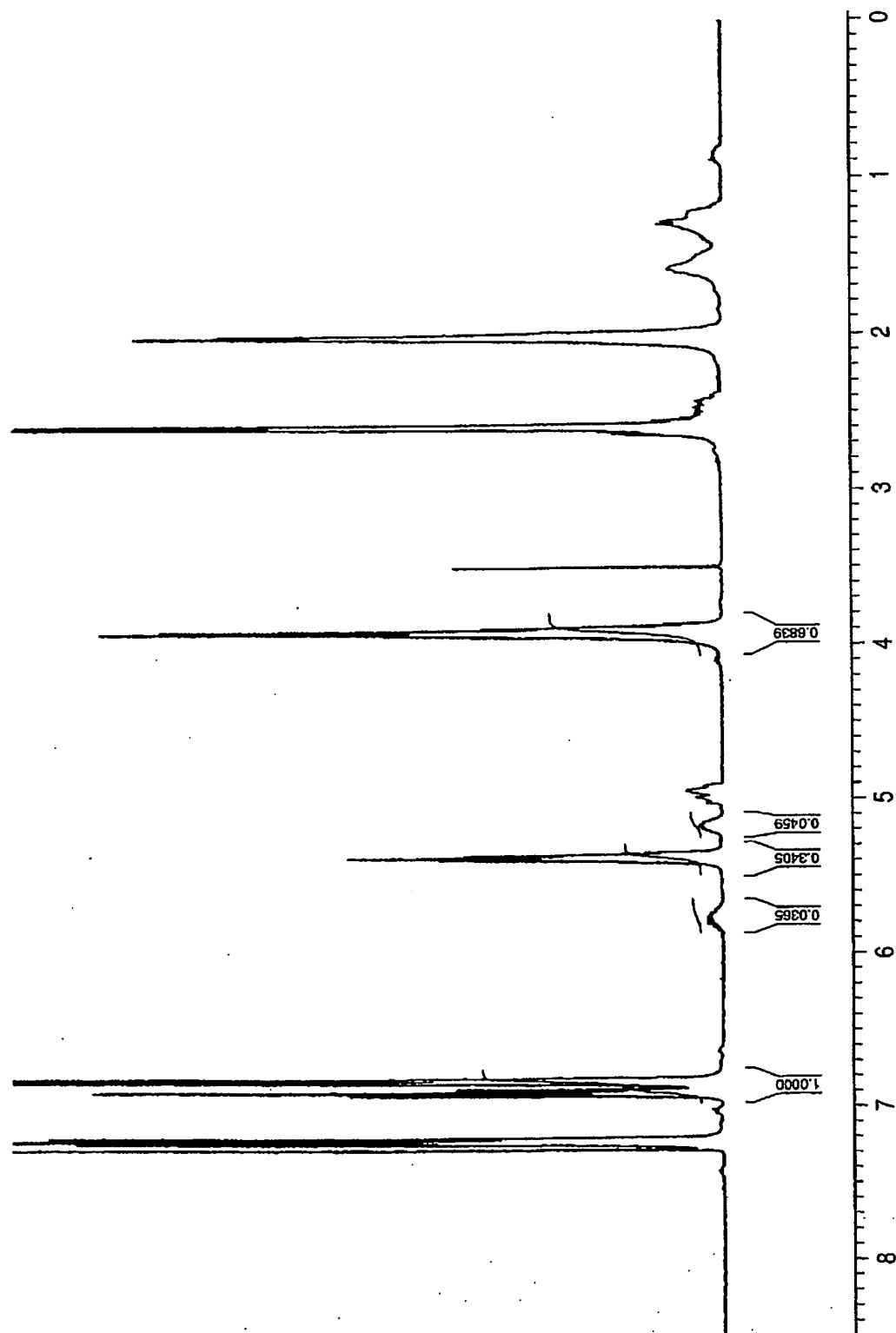
10 the chemical formula (18):



represents a (phenylmethyl)oxy group.

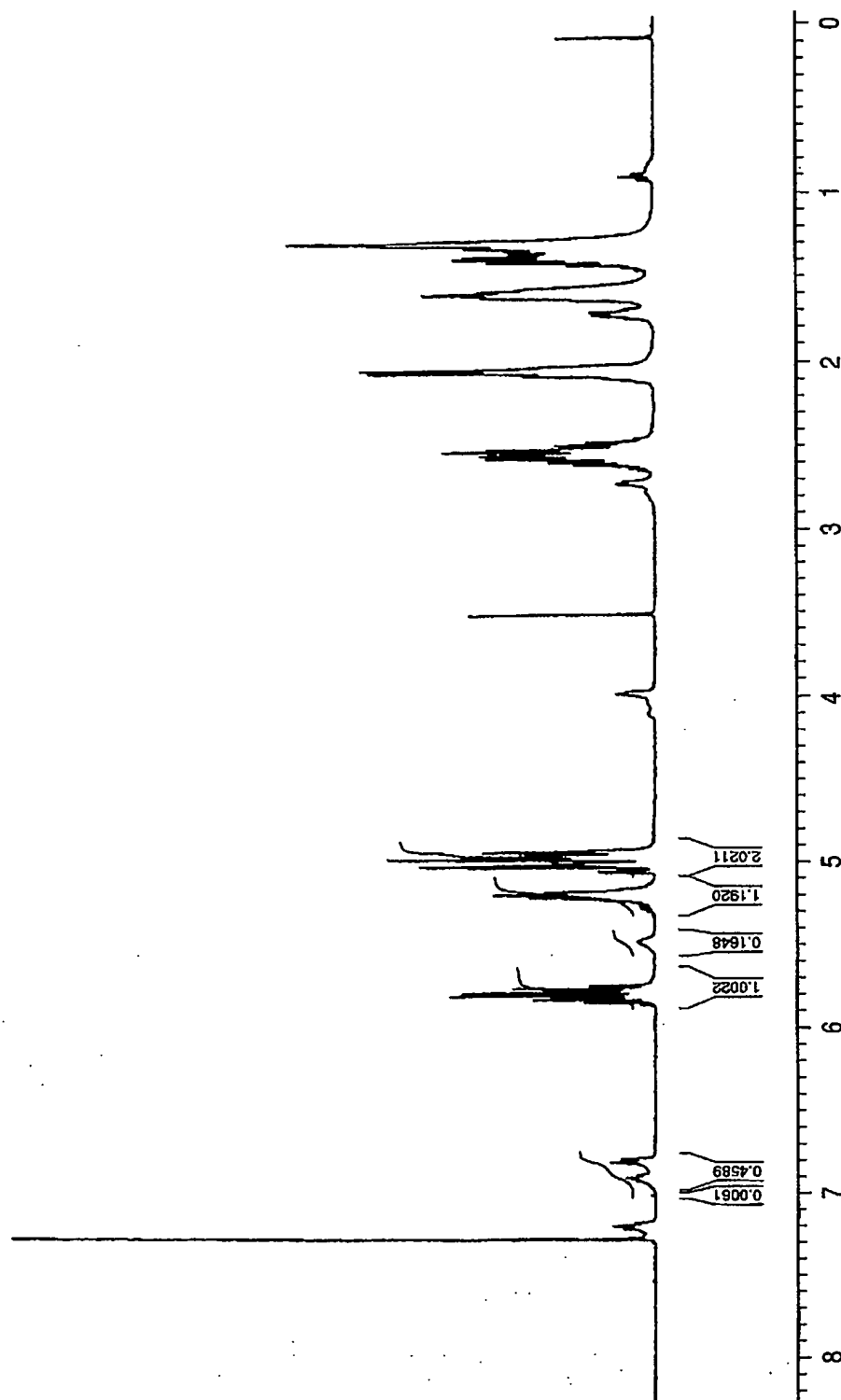
1/6

FIG. 1



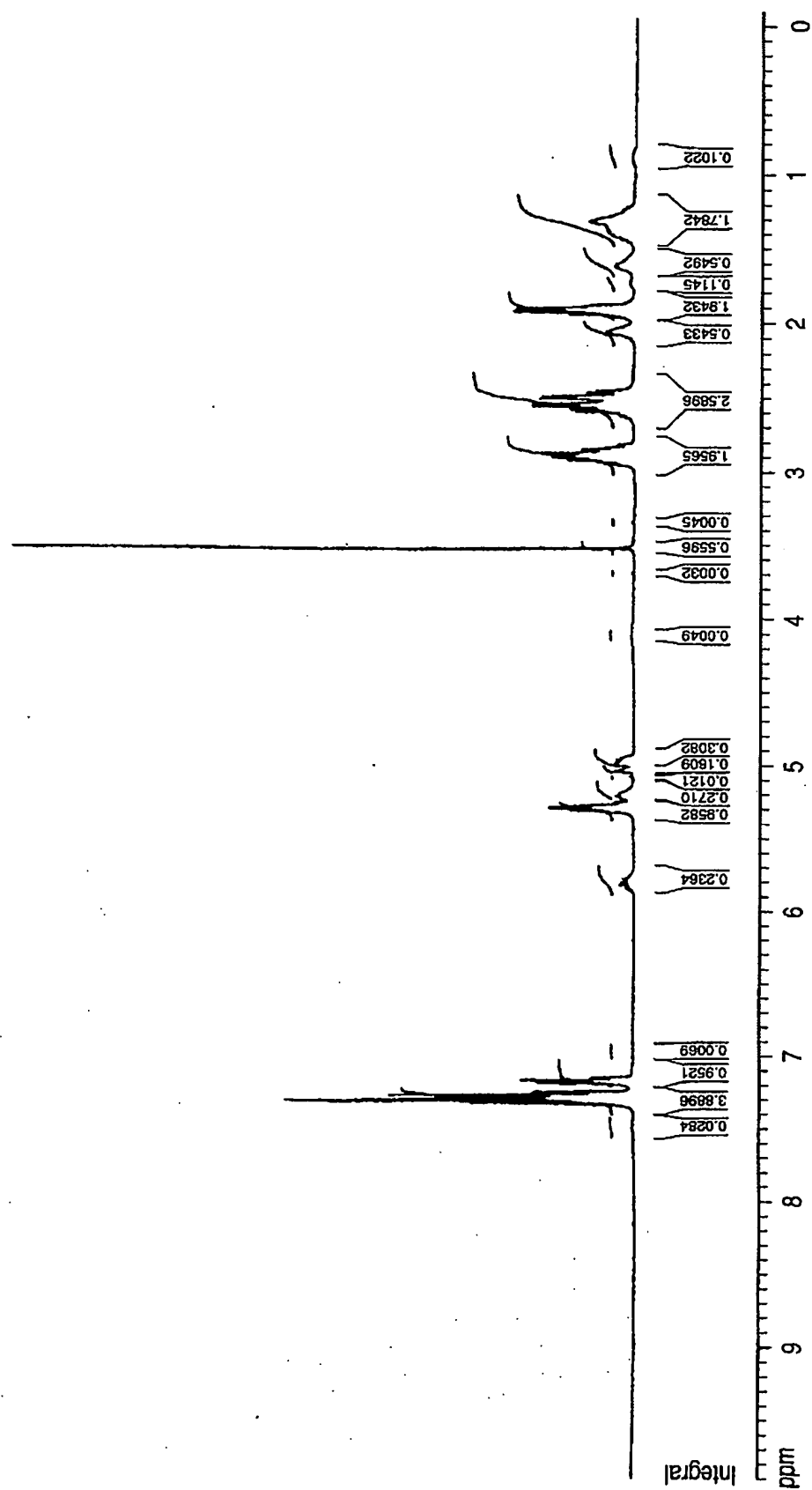
2/6

FIG. 2



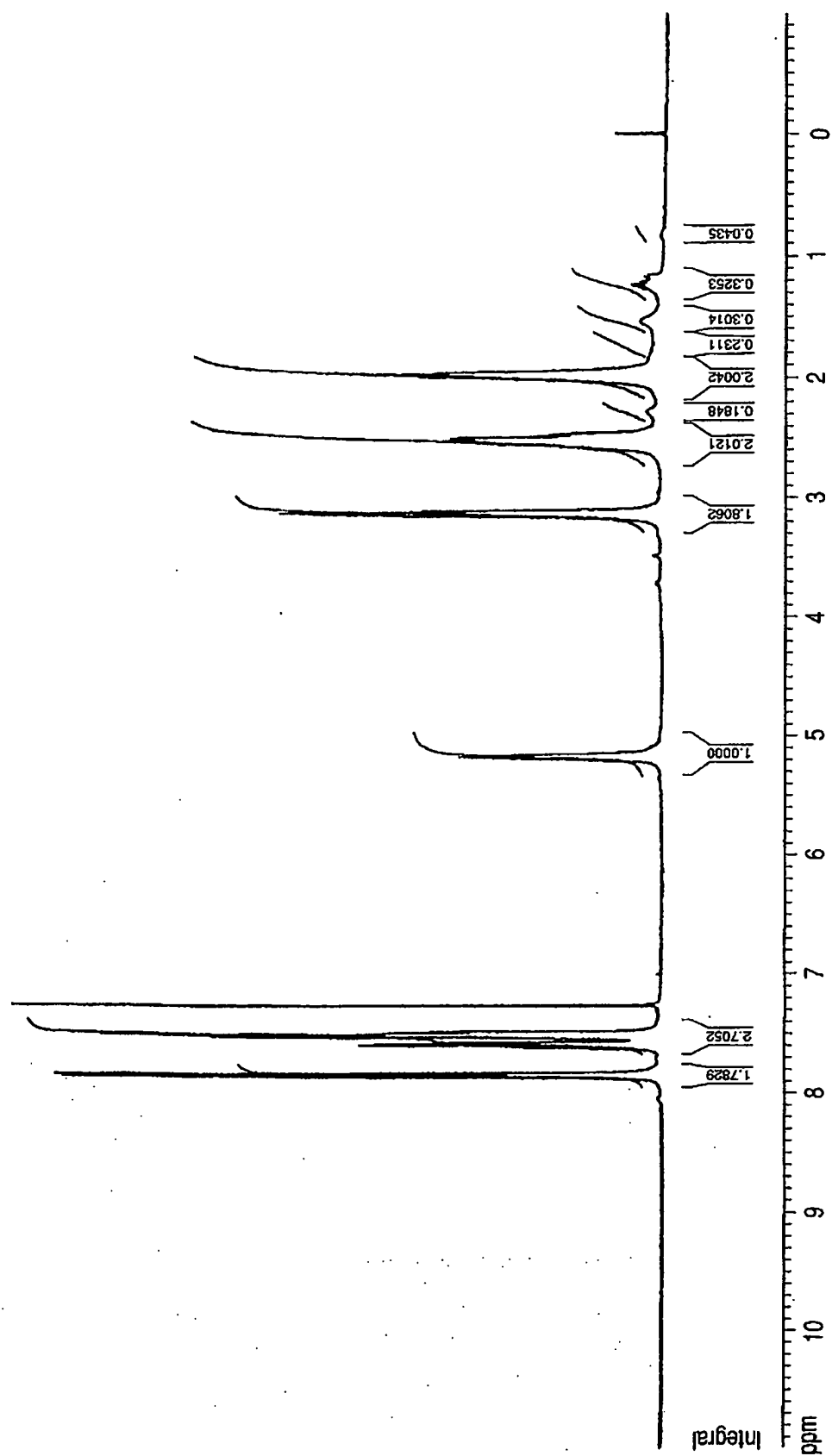
3/6

FIG. 3

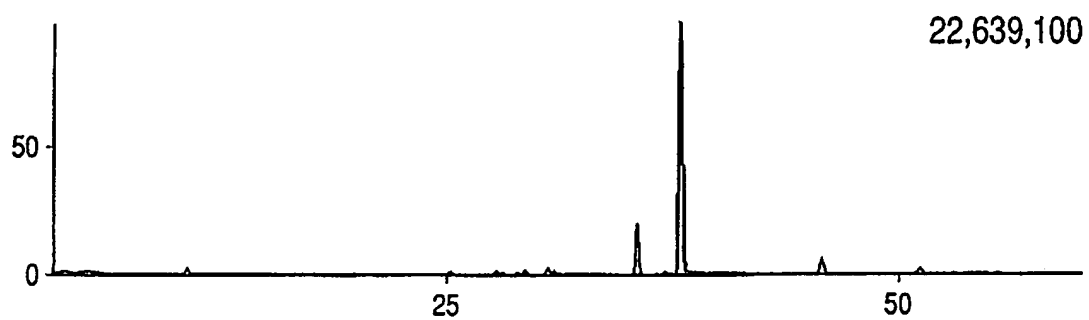
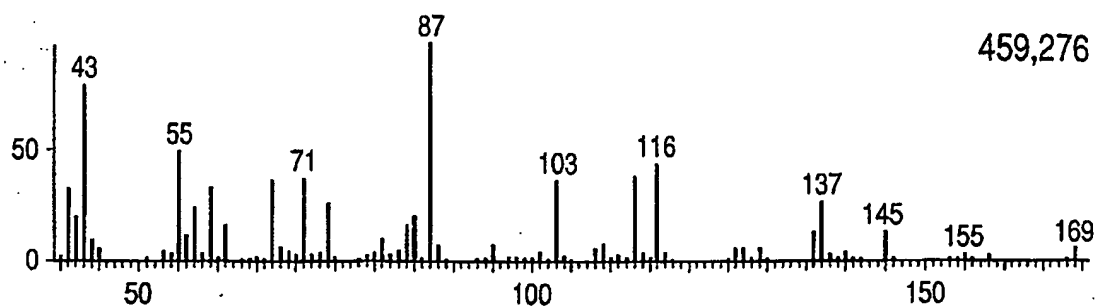


4/6

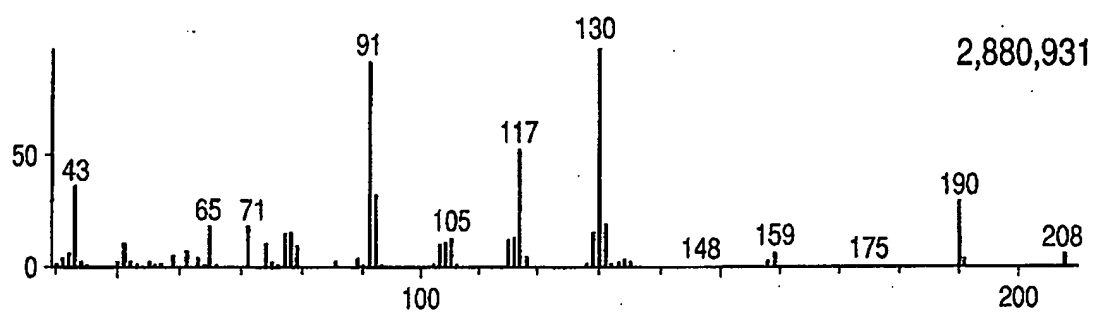
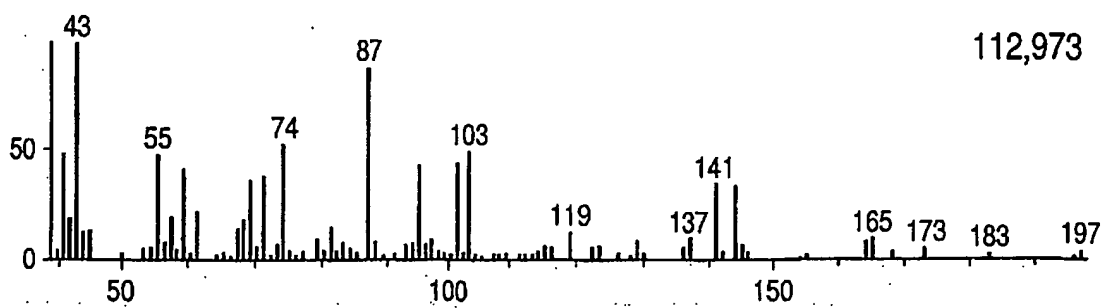
FIG. 4



5/6

FIG. 5*FIG. 6*

6/6

FIG. 7*FIG. 8*

A. CLASSIFICATION OF SUBJECT MATTER
 IPC 7 C12P7/62 C08G63/06

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
 IPC 7 C12P C08G

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, PAJ, COMPENDEX

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 5 614 576 A (RUTHERFORD DENISE R ET AL) 25 March 1997 (1997-03-25) column 5, line 19 - line 30	1-29
A	EP 1 188 836 A (CANON KK) 20 March 2002 (2002-03-20) paragraphs '0014!-'0018!; claims	1-29
A	EP 1 113 033 A (CANON KK) 4 July 2001 (2001-07-04) page 7, line 20 -page 10, line 24; claims	1-29
A	EP 1 188 782 A (CANON KK) 20 March 2002 (2002-03-20) examples	1-29
	-/-	

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents :

- 'A' document defining the general state of the art which is not considered to be of particular relevance
- 'E' earlier document but published on or after the International filing date
- 'L' document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- 'O' document referring to an oral disclosure, use, exhibition or other means
- 'P' document published prior to the International filing date but later than the priority date claimed

'T' later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

'X' document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

'Y' document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

'&' document member of the same patent family

Date of the actual completion of the international search

30 January 2004

Date of mailing of the international search report

25/02/2004

Name and mailing address of the ISA
 European Patent Office, P.B. 5818 Patentlaan 2
 NL - 2280 HV Rijswijk
 Tel (+31-70) 340-2040, Tx. 31 651 epo nl,
 Fax (+31-70) 340-3016

Authorized officer

Masson, P

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	EP 1 201 763 A (CANON KK) 2 May 2002 (2002-05-02) examples ----	1-29
A	EP 1 130 042 A (CANON KK) 5 September 2001 (2001-09-05) examples ----	1-29
A	EP 1 236 755 A (CANON KK) 4 September 2002 (2002-09-04) paragraph '0069!; claim 31; examples ----	1-29
A	EP 1 236 754 A (CANON KK) 4 September 2002 (2002-09-04) page 4, line 45 -page 7, line 4 ----	1-29
A	STEINBUECHEL A ET AL: "DIVERSITY OF BACTERIAL POLYHYDROXYALKANOIC ACIDS" FEMS MICROBIOLOGY LETTERS, AMSTERDAM, NL, vol. 128, no. 3, 15 May 1995 (1995-05-15), pages 219-228, XP000828495 ISSN: 0378-1097 page 222 -page 223; figure 2 ----	1-29
A	ANDUJAR M ET AL: "POLYESTERS PRODUCED BY PSEUDOMONAS OLEOVORANS CONTAINING CYCLOHEXYLGROUPS" MACROMOLECULES, AMERICAN CHEMICAL SOCIETY. EASTON, US, vol. 30, no. 6, 24 March 1997 (1997-03-24), pages 1611-1615, XP000682477 ISSN: 0024-9297 the whole document ----	1-29
A	EP 1 245 605 A (CANON KK) 2 October 2002 (2002-10-02) examples ----	1-29
A	SCHOLZ C., FULLER R.C. & LENZ R.W.: "Growth and Plymer Incorporation of Pseudomonas oleovorans on alkyl Esters of Heptanoic acid" MACROMOLECULES, vol. 27, 1994, pages 2886-2889, XP002268599 the whole document ----- -/--	1-29

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	<p>SCHOLZ C., FULLER R.C. & LENZ R.W.: "Production of poly(b-hydroxyalkanoates) with b-substituents containing terminal ester groups by Pseudomonas oleovorans" MACROMOLECULAR CHEMISTRY AND PHYSICS, vol. 195, 1994, pages 1405-1421, XP002268600 cited in the application the whole document</p> <p>-----</p>	1-29

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
US 5614576	A	25-03-1997	US 5753364 A	19-05-1998
			AU 700073 B2	17-12-1998
			AU 3243095 A	07-03-1996
			BR 9508583 A	14-07-1998
			CA 2195787 A1	22-02-1996
			DE 69515044 D1	16-03-2000
			DE 69515044 T2	08-06-2000
			EP 0775178 A1	28-05-1997
			ES 2141953 T3	01-04-2000
			JP 10504057 T	14-04-1998
			WO 9605264 A1	22-02-1996
EP 1188836	A	20-03-2002	JP 2002142791 A	21-05-2002
			EP 1188836 A2	20-03-2002
			US 2002052444 A1	02-05-2002
EP 1113033	A	04-07-2001	JP 2002080571 A	19-03-2002
			CN 1321695 A	14-11-2001
			EP 1113033 A2	04-07-2001
			US 2003208029 A1	06-11-2003
			US 2002022253 A1	21-02-2002
EP 1188782	A	20-03-2002	JP 2002241476 A	28-08-2002
			JP 2003047494 A	18-02-2003
			EP 1188782 A2	20-03-2002
			US 2002160467 A1	31-10-2002
EP 1201763	A	02-05-2002	JP 2002125690 A	08-05-2002
			EP 1201763 A2	02-05-2002
			US 2002081646 A1	27-06-2002
EP 1130042	A	05-09-2001	EP 1130042 A2	05-09-2001
			JP 2002173521 A	21-06-2002
			US 2003100084 A1	29-05-2003
EP 1236755	A	04-09-2002	EP 1236755 A2	04-09-2002
			JP 2003306534 A	31-10-2003
			US 2003013841 A1	16-01-2003
EP 1236754	A	04-09-2002	JP 2002327050 A	15-11-2002
			EP 1236754 A2	04-09-2002
			US 2002164726 A1	07-11-2002
EP 1245605	A	02-10-2002	EP 1245605 A2	02-10-2002
			JP 2003012786 A	15-01-2003
			US 2003096182 A1	22-05-2003